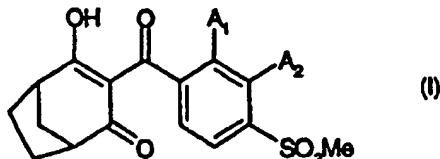




INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(51) International Patent Classification 6 : A01N 41/10		A1	(11) International Publication Number: WO 00/00029
			(43) International Publication Date: 6 January 2000 (06.01.00)
(21) International Application Number: PCT/EP99/04373			(81) Designated States: AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE, DK, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, UA, UG, US, UZ, VN, YU, ZA, ZW, ARIPO patent (GH, GM, KE, LS, MW, SD, SL, SZ, UG, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG).
(22) International Filing Date: 24 June 1999 (24.06.99)			
(30) Priority Data: 1372/98 26 June 1998 (26.06.98) CH			
(71) Applicant (for all designated States except AT US): NOVARTIS AG [CH/CH]; Schwarzwaldallee 215, CH-4058 Basel (CH).			
(71) Applicant (for AT only): NOVARTIS-ERFINDUNGEN VERWALTUNGSGESELLSCHAFT MBH [AT/AT]; Brunner Strasse 59, A-1230 Vienna (AT).			
(72) Inventor; and			Published
(75) Inventor/Applicant (for US only): RÜEGG, Willy [CH/CH]; Felmetweg 6, CH-5073 Gipf-Oberfrick (CH).			With international search report. Before the expiration of the time limit for amending the claims and to be republished in the event of the receipt of amendments.
(74) Agent: BECKER, Konrad; Novartis AG, Corporate Intellectual Property, Patent & Trademark Dept., CH-4002 Basel (CH).			

(54) Title: HERBICIDAL COMPOSITION



(57) Abstract

A herbicidal composition comprising, in addition to customary inert formulation assistants, a compound of formula (I) wherein either A₁ is nitro and A₂ is hydrogen or A₁ is methyl and A₂ is methoxy, and their salts, and b) a synergistically effective amount of one or more compounds selected from the compounds of formula (2.1) to (2.33). The compositions according to the invention may also contain a Safener.

FOR THE PURPOSES OF INFORMATION ONLY

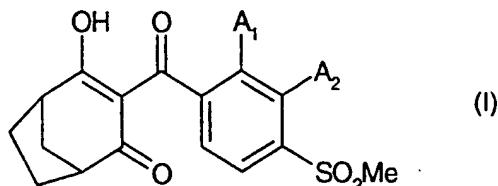
Codes used to identify States party to the PCT on the front pages of pamphlets publishing international applications under the PCT.

AL	Albania	ES	Spain	LS	Lesotho	SI	Slovenia
AM	Armenia	FI	Finland	LT	Lithuania	SK	Slovakia
AT	Austria	FR	France	LU	Luxembourg	SN	Senegal
AU	Australia	GA	Gabon	LV	Latvia	SZ	Swaziland
AZ	Azerbaijan	GB	United Kingdom	MC	Monaco	TD	Chad
BA	Bosnia and Herzegovina	GE	Georgia	MD	Republic of Moldova	TG	Togo
BB	Barbados	GH	Ghana	MG	Madagascar	TJ	Tajikistan
BE	Belgium	GN	Guinea	MK	The former Yugoslav Republic of Macedonia	TM	Turkmenistan
BF	Burkina Faso	GR	Greece	ML	Mali	TR	Turkey
BG	Bulgaria	HU	Hungary	MN	Mongolia	TT	Trinidad and Tobago
BJ	Benin	IE	Ireland	MR	Mauritania	UA	Ukraine
BR	Brazil	IL	Israel	MW	Malawi	UG	Uganda
BY	Belarus	IS	Iceland	MX	Mexico	US	United States of America
CA	Canada	IT	Italy	NE	Niger	UZ	Uzbekistan
CF	Central African Republic	JP	Japan	NL	Netherlands	VN	Viet Nam
CG	Congo	KE	Kenya	NO	Norway	YU	Yugoslavia
CH	Switzerland	KG	Kyrgyzstan	NZ	New Zealand	ZW	Zimbabwe
CI	Côte d'Ivoire	KP	Democratic People's Republic of Korea	PL	Poland		
CM	Cameroon	KR	Republic of Korea	PT	Portugal		
CN	China	KZ	Kazakhstan	RO	Romania		
CU	Cuba	LC	Saint Lucia	RU	Russian Federation		
CZ	Czech Republic	LI	Liechtenstein	SD	Sudan		
DE	Germany	LK	Sri Lanka	SE	Sweden		
DK	Denmark	LR	Liberia	SG	Singapore		
EE	Estonia						

Herbicidal composition

The present invention relates to a novel herbicidal synergistic composition that contains a combination of herbicides suitable for selectively controlling weeds in crops of cultivated plants, typically in crops of maize. The invention further relates to a process for controlling weeds in crops of cultivated plants and to the use of said novel composition for this purpose.

Compounds of formula I

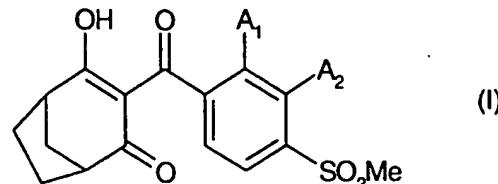


wherein either A₁ is nitro and A₂ is hydrogen or A₁ is methyl and A₂ is methoxy, and their salts, possess herbicidal activity, as is described for example in EP-A-0 338 992.

It has now surprisingly been shown that a combination of active compounds, in a ratio varying within specific limits, i.e. an active ingredient of formula I with one or more of the herbicides of formulae 2.1 to 2.33 listed below, which are known and are partly available commercially, exerts a synergistic effect that is able to control the majority of weeds occurring preferably in crops of cultivated plants preemergence as well as postemergence.

Accordingly, the present invention provides a novel synergistic composition for the selective control of weeds that, in addition to customary inert formulation assistants, contains as active ingredient a mixture of

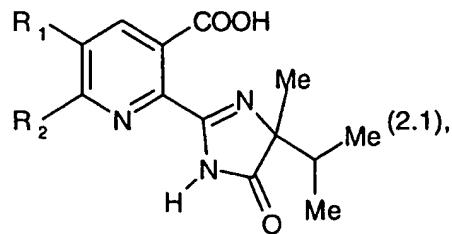
a) a compound of formula I



wherein either A₁ is nitro and A₂ is hydrogen or A₁ is methyl and A₂ is methoxy, as well as their salts, and

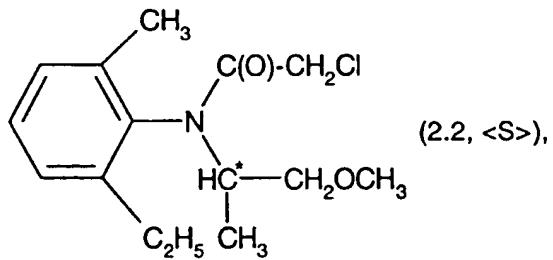
- 2 -

b) a synergistically active amount of one or more compounds selected from the compound of formula 2.1

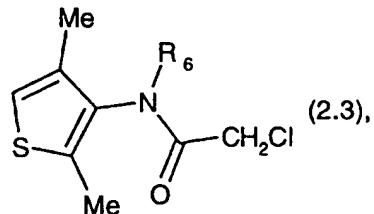


wherein R₁ is CH₂-OMe, ethyl or hydrogen,

R₂ is hydrogen or R₁ and R₂ together are the group -CH=CH-CH=CH-; and the compound of formula 2.2

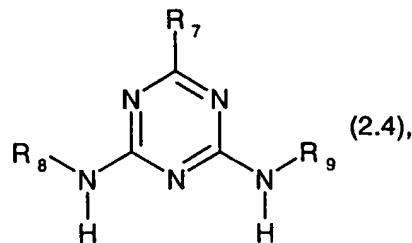


and the compound of formula 2.3



wherein R₆ is CH(Me)-CH₂OMe or <S>CH(Me)-CH₂OMe;

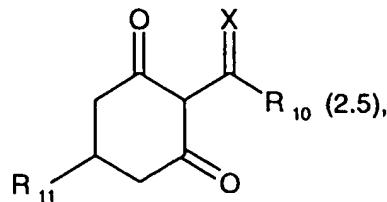
and the compound of formula 2.4



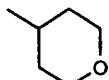
wherein R₇ is chlorine or SMe, R₈ is ethyl and R₉ is ethyl, isopropyl or tert.-butyl;

and the compound of formula 2.5

- 3 -

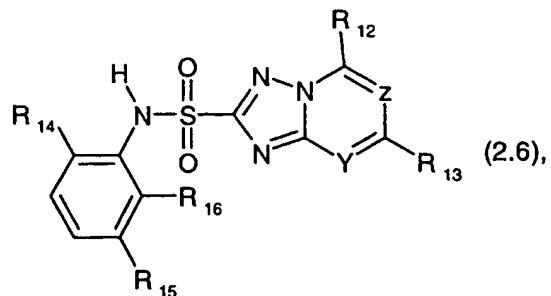


wherein R_{10} is ethyl or n-propyl, R_{11} is $\text{COO}^- \text{ 1/2 Ca}^{++}$, $-\text{CH}_2\text{CH}(\text{Me})\text{S}-\text{CH}_2\text{CH}_3$ or the group



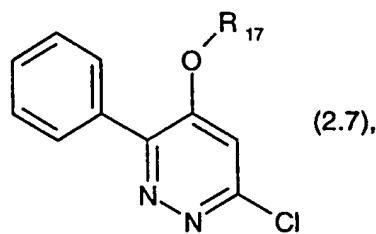
and X is oxygen, $\text{N}-\text{O}-\text{CH}_2\text{CH}_3$ or $\text{N}-\text{O}-\text{CH}_2\text{CH}=\text{CH}-\text{Cl}$;

and the compound of formula 2.6



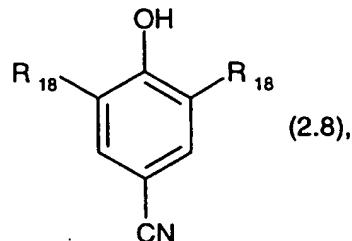
wherein R_{12} is hydrogen, methoxy or ethoxy, R_{13} is Me, methoxy or fluorine, R_{14} is COOMe , fluorine or chlorine, R_{15} is hydrogen or Me, Y is methine or nitrogen, Z is methine or nitrogen and R_{16} is fluorine or chlorine;

and the compound of formula 2.7



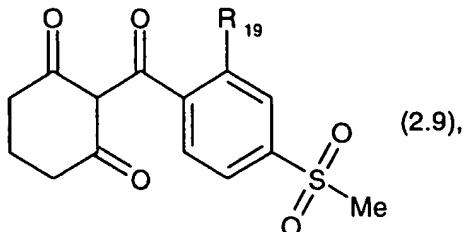
wherein R_{17} is hydrogen or $-\text{C}(\text{O})-\text{S}-\text{n-octyl}$;

and the compound of formula 2.8

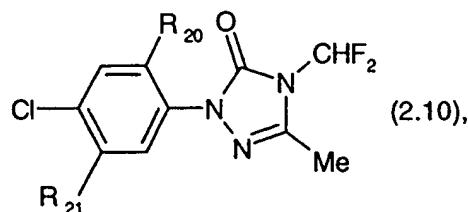


- 4 -

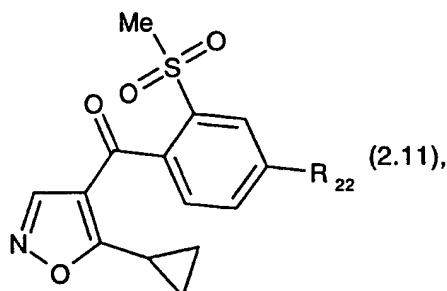
when R_{18} is bromine or iodine;
and the compound of formula 2.9



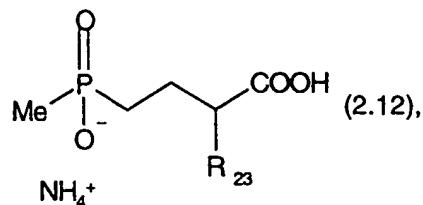
wherein R_{19} is chlorine or nitro;
and the compound of formula 2.10



wherein R_{20} is fluorine or chlorine and R_{21} is $-\text{CH}_2\text{-CH}(\text{Cl})\text{-COOCH}_2\text{CH}_3$ or $-\text{NH-SO}_2\text{Me}$; and the compound of formula 2.11

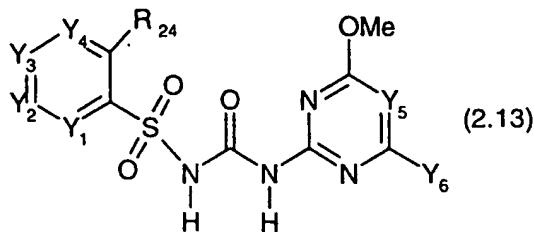


wherein R₂₂ is trifluoromethyl or chlorine; and the compound of formula 2.12



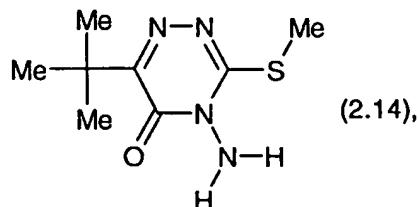
wherein R₂₃ is NH₂ or <S>NH₂;
and the compound of formula 2.13

- 5 -

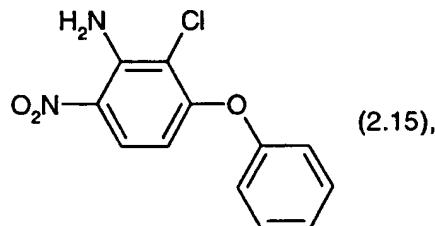


wherein Y_1 is nitrogen, methine or $N\text{-Me}$, Y_2 is nitrogen, methine or $C\text{-I}$, Y_3 is methine, Y_4 is methine, or Y_3 and Y_4 together are sulphur or $C\text{-Cl}$, Y_5 is nitrogen or methine, Y_6 is methyl or methoxy and R_{24} is CONMe_2 , COOMe , $\text{CH}_2\text{-CH}_2\text{CF}_3$ or $\text{SO}_2\text{CH}_2\text{CH}_3$, or the sodium salts thereof;

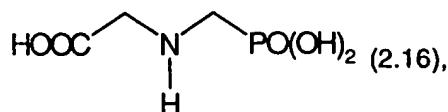
and the compound of formula 2.14



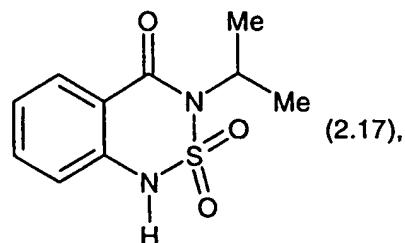
and the compound of formula 2.15



and the compound of formula 2.16

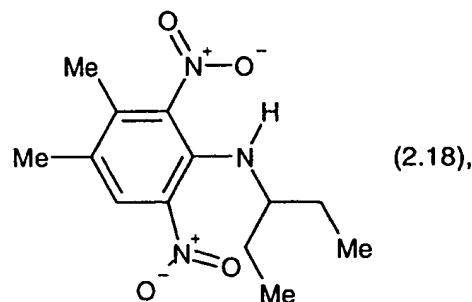


and the compound of formula 2.17

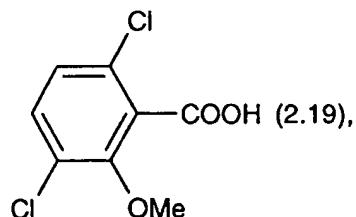


- 6 -

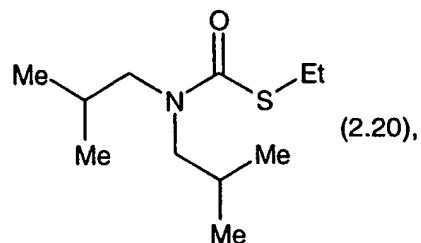
and the compound of formula 2.18



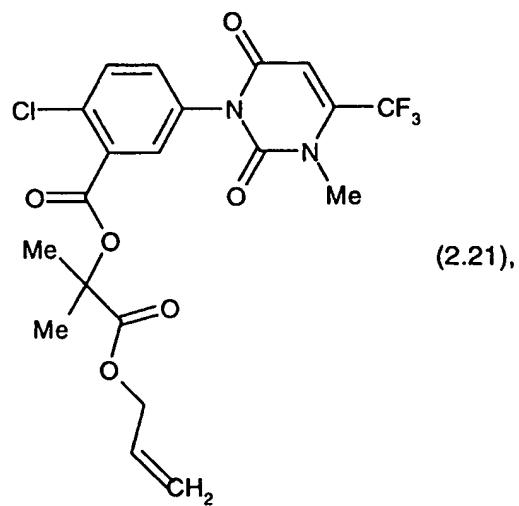
and the compound of formula 2.19



and the compound of formula 2.20

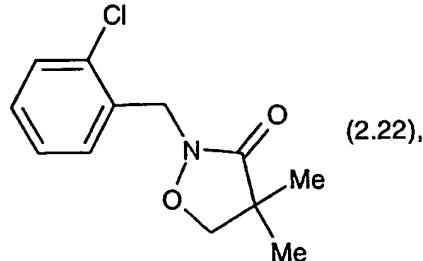


and the compound of formula 2.21

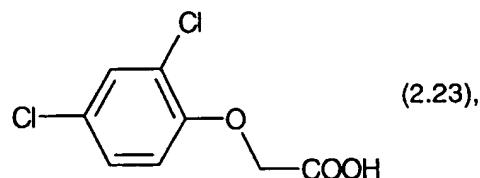


- 7 -

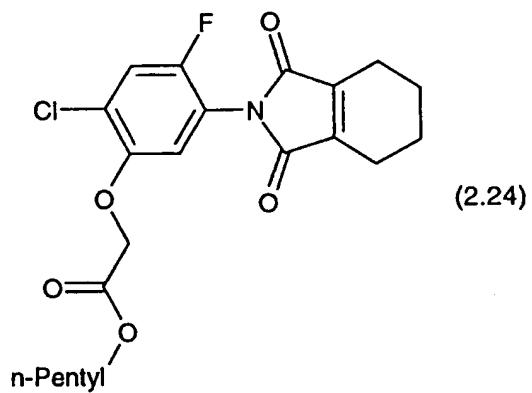
and the compound of formula 2.22



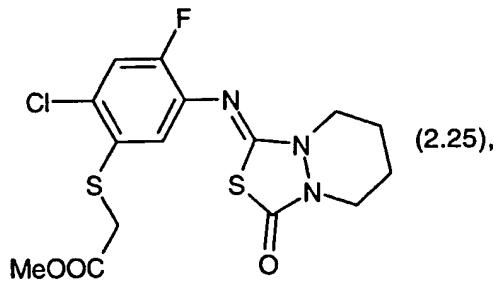
and the compound of formula 2.23



and the compound of formula 2.24

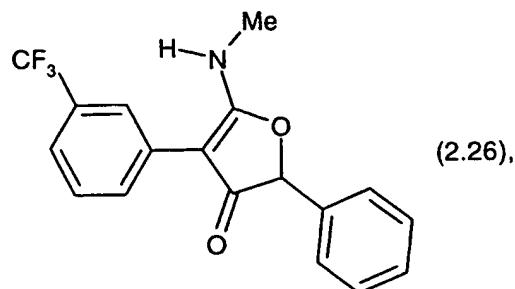


and the compound of formula 2.25

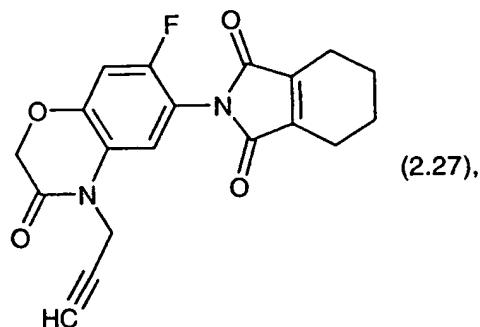


and the compound of formula 2.26

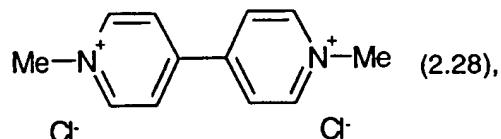
- 8 -



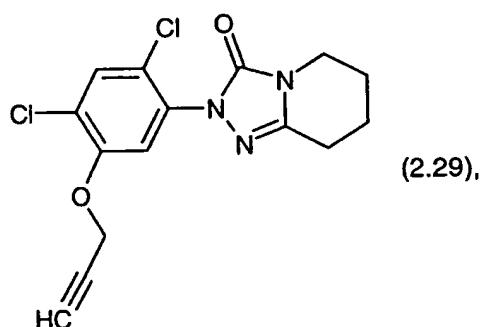
and the compound of formula 2.27



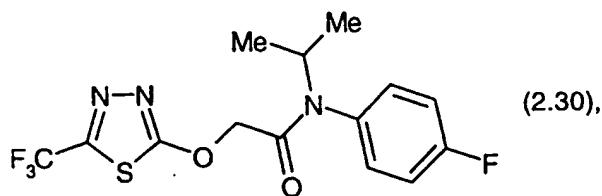
and the compound of formula 2.28



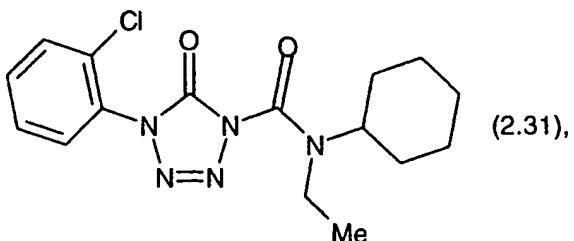
and the compound of formula 2.29



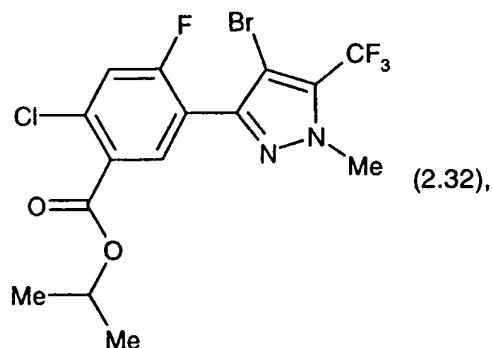
and the compound of formula 2.30



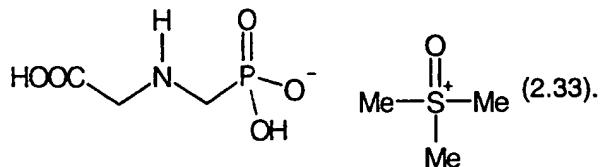
and the compound of formula 2.31



and the compound of formula 2.32



and the compound of formula 2.33



In the above formulae, "Me" signifies the methyl group. The invention also embraces the salts that the compounds of formula I are able to form with amines, alkali metal bases and alkaline earth metal bases or quarternary ammonium bases. Salt-forming alkali metal and alkaline earth metal hydroxides include the hydroxides of lithium, sodium, potassium, magnesium or calcium, those of sodium or potassium being especially preferred.

Illustrative examples of amines suitable for forming ammonium salts are ammonia, as well as primary, secondary, and tertiary C₁-C₁₈alkylamines, C₁-C₄hydroxyalkylamines and C₂-C₄alkoxyalkylamines, typically methylamine, ethylamine, n-propylamine, isopropylamine, the four isomeric butylamines, n-amylamine, isoamylamine, hexylamine, heptylamine, octylamine, nonylamine, decylamine, pentadecylamine, hexadecylamine, heptadecylamine, octadecylamine, methyl ethylamine, methyl isopropylamine, methyl hexylamine, methyl

nonylamine, methyl pentadecylamine, methyl octadecylamine, ethyl butylamine, ethyl heptylamine, ethyl octylamine, hexyl heptylamine, hexyl octylamine, dimethylamine, diethylamine, di-n-propylamine, diisopropylamine, di-n-butylamine, di-n-amylamine, diisoamylamine, dihexylamine, diheptylamine, dioctylamine, ethanolamine, n-propanolamine, isopropanolamine, N,N-diethanolamine, N-ethylpropanolamine, N-butylethanolamine, allylamine, n-butenyl-2-amine, n-pentenyl-2-amine, 2,3-dimethylbutenyl-2-amine, dibutenyl-2-amine, n-hexenyl-2-amine, propylenediamine, trimethylamine, triethylamine, tri-n-propylamine, triisopropylamine, tri-n-butylamine, triisobutylamine, tri-sec-butylamine, tri-n-amylamine, methoxyethylamine and ethoxyethylamine; heterocyclic amines such as pyridine, quinoline, isoquinoline, morpholine, piperidine, pyrrolidine, indoline, quinuclidine and azepine; primary arylamines such as anilines, methoxyanilines, ethoxyanilines, o-, m- and p-toluidines, phenylenediamines, benzidines, naphthylamines and o-, m- and p-chloroanilines. Preferred amines are triethylamine, isopropylamine and diisopropylamine. The compound of formula I, wherein A₁ is methyl and A₂ is methoxy, is most preferably present in the form of the diisopropylammonium salt.

It is entirely surprising that the combination of the compound of formula I with one or more herbicides selected from formulae 2.1 to 2.33 exceeds the expected additive action against the weeds to be controlled and thus in particular enhances the activity range of the individual components in two respects: On the one hand, the concentrations of the single compounds of formulae I and 2.1 to 2.33 are reduced whilst retaining good activity. On the other hand, the novel herbicidal combination also achieves a high degree of weed control where the single compounds have become no longer agriculturally effective at low concentrations. The consequence is a substantial broadening of the activity spectrum against weeds and an additional increase in the selectivity for the cultivated plants that is necessary and desirable in the event of unintentional overapplication of herbicide. In addition, the novel composition permits greater flexibility with respect to subsequent crops while retaining the excellent control of weeds in crops of cultivated plants.

The composition of the invention may be used against a large number of agronomically important weeds, typically *Stellaria*, *Nasturtium*, *Agrostis*, *Digitaria*, *Avena*, *Setaria*, *Sinapis*, *Lolium*, *Solanum*, *Phaseolus*, *Echinochloa*, *Scirpus*, *Monochoria*, *Sagittaria*, *Bromus*, *Alopecurus*, *Sorghum halepense*, *Rottboellia*, *Cyperus*, *Abutilon*, *Sida*, *Xanthium*, *Amaranthus*, *Chenopodium*, *Ipomoea*, *Chrysanthemum*, *Galium*, *Viola*, and *Veronica*. The

composition of this invention is suitable for all methods of application commonly used in agriculture, including preemergence application, postemergence application and seed dressing. The composition of the invention is preferably suitable for weed control in crops of cultivated plants, typically cereals, rape, sugar beet, sugar cane, plantations, rice, maize and soybeans and for the non-selective control of weeds.

Crops will also be understood as meaning those crops that have been made tolerant to herbicides or classes of herbicides by conventional breeding or genetic engineering methods.

The compounds of formulae 2.1 and 2.3 to 2.13 are known under the names imazamox, imazethapyr, imazaquin, imazapyr, dimethenamid, atrazine, terbutylazin, simazine, terbutyln, prohexadione calcium, sethoxydim, clethodim, tepraloxymid, flumetsulam, metosulam, pyridate, bromoxynil, ioxynil, sulcotrione, carfentrazone, sulfentrazone, isoxaflutole, glufosinate, primisulfuron, prosulfuron, rimsulfuron, halosulfuron, nicosulfuron, and thifensulfuron and are described in the Pesticide Manual, eleventh ed., British Crop Protection Council, 1997 under the entry numbers 412, 415, 414, 413, 240, 34, 692, 651, 693, 595, 648, 146, 49, 339, 495, 626, 88, 425, 664, 112, 665, 436, 382, 589, 613, 644, 389, 519 and 704. The compound of formula 2.13, wherein Y_1 , Y_3 and Y_4 are methine, Y_2 is C-I, R_{24} is COOMe, Y_5 is nitrogen and Y_6 is methyl, is known under the name iodosulfuron (especially the sodium salt) from AGROW No. 296, 16th January 1998, page 22. The S-enantiomer of the compound of formula 2.12 is registered under CAS reg. no. [35597-44-5]. The compound of formula 2.2, aRS,1'S(-)-N-(1'-methyl-2'-methoxyethyl)-N-chloroacetyl-2-ethyl-6-methylaniline, as well as a compound of the general formula 2.3, (1S,aRS)-2-chloro-N-(2,4-dimethyl-3-thienyl)-N-(2-methoxy-1-methylethyl)-acetamide, are described for example in WO 97/34485. The compound of formula 2.9, wherein R_{19} is NO₂, is known under the name mesotrione and is described for example in US-A-5.006.158. The compound of formula 2.6, wherein R_{12} is ethoxy, R_{13} is fluorine, Y is methine, R_{14} is methoxycarbonyl, R_{15} is hydrogen and R_{16} is chlorine, is known under the name cloransulam, for example from AGROW no. 261, 2nd August 1996, page 21.

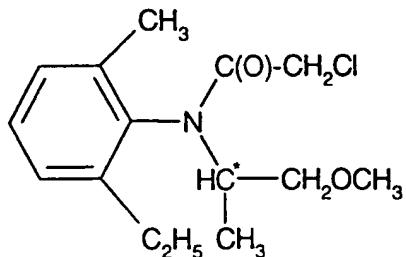
In addition, the following compounds of the composition according to the invention are described in the Pesticide Manual, eleventh ed., British Crop Protection Council, 1997:

Compound of formula (name)	Pesticide Manual eleventh ed., Entry No.:
2.14 (Metribuzin)	497
2.15 (Aclonifen)	8
2.16 (Glyphosate)	383
2.17 (Bentazone)	65
2.18 (Pendimethalin)	557
2.19 (Dicamba)	210
2.20 (Butylate)	100
2.22 (Clomazone)	150
2.23 (2,4-D)	192
2.24 (Flumiclorac)	340
2.25 (Fluthiacet-methyl)	359
2.26 (Flurtamone)	356
2.27 (Flumioxazin)	341
2.28 (Paraquat)	550
2.29 (Azafenidin)	37
2.30 (Fluthiamid)	51
2.33 (Sulfosate)	383

The compound of formula 2.7, wherein R₁₇ is hydrogen, and the preparation thereof, are described in US-A-3,790,571, the compound of formula 2.6, wherein R₁₂ is ethoxy, Z is nitrogen, R₁₃ is fluorine, R₁₄ is chlorine and R₁₅ is hydrogen and R₁₆ is chlorine, is described in US-A-5,498,773.

The compound of formula 2.21 and the preparation thereof are described in US-A-5,183,492, the compound of formula 2.22 is described under the name isoxachlortole in AGROW no. 296, 16th January 1998, page 22. The compound of formula 2.31 is described under the name fentrazamide in The 1997 British Crop Protection Conference - Weeds, Conference Proceedings Vol. 1, 2-8, pages 67 to 72, the compound of formula 2.32 is described under the name JV 485 (Isoxapropazol) in The 1997 British Crop Protection Conference - Weeds, Conference Proceedings Vol. 1, 3A-2, pages 93 to 98.

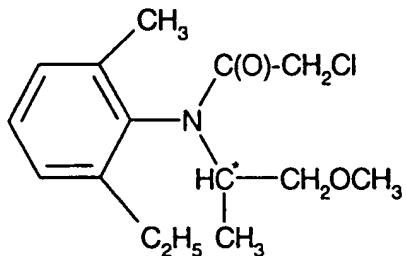
Preferred synergistic mixtures according to the invention contain as active ingredients a compound of formula I and either the compound of formula 2.2



(2,2, aRS,1'S(-)N-(1'-methyl-2'-methoxyethyl)-N-

chloroacetyl-2-ethyl-6-methylaniline), or a compound of formula 2.3, or a compound of formula 2.4, wherein R₇ is chlorine, R₈ is ethyl and R₉ is isopropyl, or R₇ is chlorine, R₈ is ethyl and R₉ is tert.-butyl, or a compound of formula 2.6, wherein R₁₂ is hydrogen, Z is methine, R₁₃ is methyl, Y is nitrogen, R₁₄ is fluorine, R₁₅ is hydrogen and R₁₆ is fluorine, or R₁₂ is methoxy, Z is methine, R₁₃ is methoxy, Y is methine, R₁₄ is chlorine, R₁₅ is methyl and R₁₆ is chlorine, or a compound of formula 2.7, wherein R₁₇ is $-\text{C}(\text{O})\text{-S-n-octyl}$, or a compound of formula 2.9, or a compound of formula 2.11, wherein R₂₂ is trifluoromethyl, or a compound of formula 2.12, or a compound of formula 2.13, wherein Y₁ is nitrogen, Y₂, Y₃ and Y₄ are methine, R₂₄ is dimethylaminocarbonyl and Y₅ is methine, or Y₁, Y₂, Y₃ and Y₄ are methine, R₂₄ is methoxycarbonyl and Y₅ is methine, or Y₁, Y₂, Y₃ and Y₄ are methine, R₂₄ is 3-trifluoropropyl and Y₅ is nitrogen, or Y₁ is nitrogen, Y₂, Y₃ and Y₄ are methine, R₂₄ is ethylsulphonyl and Y₅ is methine, or Y₁ is N-Me, Y₂ is nitrogen, Y₃ and Y₄ together are C-Cl, R₂₄ is methoxycarbonyl and Y₅ is methine, or a compound of formula 2.16, or a compound of formula 2.18, or a compound of formula 2.19, or a compound of formula 2.30.

A further group of preferred synergistic mixtures according to the invention contains as active ingredients a compound of formula I, the compound of formula 2.2

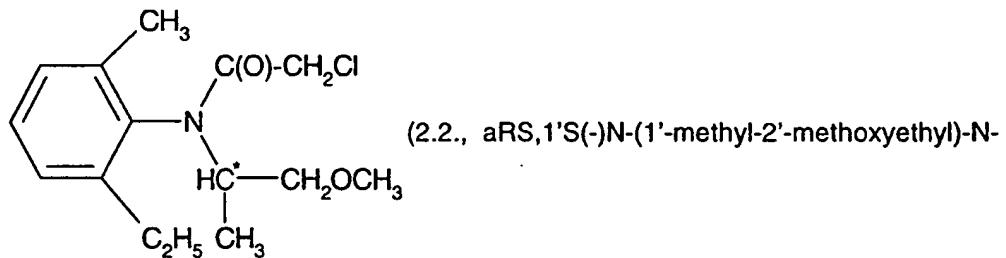


(2,2, aRS,1'S(-)N-(1'-methyl-2'-methoxyethyl)-N-

chloroacetyl-2-ethyl-6-methylaniline), as well as a compound selected from formula 2.4, wherein R₇ is chlorine, R₈ is ethyl and R₉ is isopropyl, or R₇ is chlorine, R₈ is ethyl and R₉ is tert.-butyl, and of formula 2.6, wherein R₁₂ is hydrogen, Z is methine, R₁₃ is methyl, Y is

nitrogen, R₁₄ is fluorine, R₁₅ is hydrogen and R₁₆ is fluorine, or R₁₂ is methoxy, Z is methine, R₁₃ is methoxy, Y is methine, R₁₄ is chlorine, R₁₅ is methyl and R₁₆ is chlorine, and of formula 2.7, wherein R₁₇ is -C(O)-S-n-octyl, and of formula 2.13, wherein Y₁ is nitrogen, Y₂, Y₃ and Y₄ are methine, R₂₄ is dimethylaminocarbonyl and Y₅ is methine, or Y₁, Y₂, Y₃ and Y₄ are methine, R₂₄ is methoxycarbonyl and Y₅ is methine, or Y₁, Y₂, Y₃ and Y₄ are methine, R₂₄ is 3-trifluoropropyl and Y₅ is nitrogen, or Y₁ is nitrogen, Y₂, Y₃ and Y₄ are methine, R₂₄ is ethylsulphonyl and Y₅ is methine, or Y₁ is N-Me, Y₂ is nitrogen, Y₃ and Y₄ together are C-Cl, R₂₄ is methoxycarbonyl and Y₅ is methine, and of formula 2.9, and of formula 2.11, wherein R₂₂ is trifluoromethyl, and of formula 2.12, and of formula 2.16, and of formula 2.18, and of formula 2.19.

It has been found that particularly effective herbicidal combinations are the combinations of compounds of formula I with the compound of formula 2.2



The composition according to the invention contains the active ingredient of formula I and the active ingredients of formulae 2.1 to 2.33 in any ratio, normally with an excess of one component over the others. In general, the mixture ratios (weight ratios) between the active ingredient of formula I and the components of formulae 2.1 to 2.33 lie between 1:2000 and 2000:1, especially between 200:1 and 1:200.

The rate of application can vary over a wide range and will depend on the nature of the soil, the type of application (pre- or postemergence), seed dressing, application to the seed furrow; no tillage application etc.), the cultivated plant, the weed to be controlled, the respective prevailing climatic conditions; and on other factors governed by the type and time of application and the target crop. In general, the mixture according to the invention may be used at a rate of application of 1 to 5000 g of mixture/ha.

The mixtures of the compounds of formula I with the compounds of formulae 2.1 to 2.33 may be used in unmodified form, i.e. as obtained in the synthesis. Preferably, however, they are processed in conventional manner e.g. to emulsifiable concentrates, directly sprayable or dilutable solutions, dilute emulsions, wettable powders, soluble powders, dusts, granulates or microcapsules using the auxiliary agents customarily employed in formulation technology. As with the type of compositions, the methods of application – such as spraying, atomising, dusting, wetting, scattering, or pouring – are selected in accordance with the intended objectives and the prevailing circumstances.

The formulations, i.e. the agents, preparations, or compositions containing the compounds of formula I and 2.1 to 2.33, and optionally one or more than one liquid or solid formulation assistant, are prepared in known manner, e.g. by homogeneously mixing and/or grinding the herbicide with said formulation auxiliaries, typically solvents or solid carriers. Surface-active compounds (surfactants) may additionally be used for preparing the formulations.

Examples of solvents and solid carriers are described in WO 97/34485, page 6.

Depending on the herbicide of formula I to be formulated, suitable surface-active compounds are nonionic, cationic and/or anionic surfactants and surfactant mixtures having good emulsifying, dispersing and wetting properties.

Examples of suitable anionic, non-ionic, and cationic surfactants are listed in WO 97/34485 on pages 7 and 8.

Also the surfactants customarily for the art of formulation and described, *inter alia*, in "Mc Cutcheon's Detergents and Emulsifiers Annual" MC Publishing Corp., Ridgewood New Jersey, 1981, Stache, H., "Tensid-Taschenbuch" (Handbook of Surfactants), Carl Hanser Verlag, Munich/Vienna, 1981, and M. and J. Ash, "Encyclopedia of Surfactants", Vol I-III, Chemical Publishing Co., New York, 1980-81 are suitable for manufacture of the herbicides according to the invention.

The herbicidal compositions will usually contain from 0.1 to 99% by weight, preferably from 0.1 to 95% by weight, of compound mixture of the compound of formula I and the compounds of formulae 2.1 to 2.33, from 1 to 99.9% by weight of a solid or liquid formulation assistant, and from 0 to 25% by weight, preferably from 0.1 to 25% by weight, of a surfactant.

Whereas it is customarily preferred to formulate commercial products as concentrates, the end user will normally use dilute formulations. The compositions may also contain further ingredients, such as: stabilisers, e.g. where appropriate epoxidised vegetable oils (epoxidised coconut oil, rapeseed oil, or soybean oil); antifoams, typically silicone oil; preservatives; viscosity regulators; binders; and tackifiers; as well as fertilisers or other chemical agents. Particularly preferred formulations are made up as follows:

(% = percent by weight)

Emulsifiable concentrates:

Compound mixture: 1 to 90 %, preferably 5 to 20%
Surfactant: 1 to 30%, preferably 10 to 20%
Liquid carrier: 5 to 94 %, preferably 70 to 85 %

Dusts:

Compound mixture: 0.1 to 10 %, preferably 0.1 to 5 %
Solid carrier: 99.9 to 90 %, preferably 99.9 to 99 %

Suspension concentrates:

Compound mixture: 5 to 75 %, preferably 10 to 50 %
Water: 94 to 24%, preferably 88 to 30%
Surfactant: 1 to 40 %, preferably 2 to 30%

Wettable powders:

Compound mixture: 0.5 to 90 %, preferably 1 to 80 %
Surfactant: 0.5 to 20 %, preferably 1 to 15 %
Solid carrier: 5 to 95 %, preferably 15 to 90 %

Granulates:

Compound mixture: 0.1 to 30 %, preferably 0.1 to 15 %

Solid carrier: 99.5 to 70 %, preferably 97 to 85 %

The invention is illustrated by the following non-limitative Examples.

F1. Emulsifiable concentrates

	a)	b)	c)	d)
Compound mixture	5 %	10 %	25 %	50 %
Calcium dodecylbenzenesulfonate	6 %	8 %	6 %	8 %
Polyethoxylated castor oil (36 mols EO)	4 %	-	4 %	4 %
Octylphenol polyethoxylate (7-8 mols EO)	-	4 %	-	2 %
Cyclohexanone	-	-	10 %	20 %
Aromatic hydrocarbon mixture C ₉ -C ₁₂	85 %	78 %	55 %	16 %

Emulsions of any desired concentration can be prepared by diluting such concentrates with water.

F2. Solutions

	a)	b)	c)	d)
Compound mixture	5 %	10 %	50 %	90 %
1-methoxy-3-(3-methoxy- propoxy)-propane	-	20 %	20 %	-
Polyethylene glycol mw 400	20 %	10 %	-	-
N-Methyl-2-pyrrolidone	-	-	30 %	10 %
Aromatic hydrocarbon mixture C ₉ -C ₁₂	75 %	60 %	-	-

The solutions are suitable for use in the form of microdrops.

F3. Wettable powders

	a)	b)	c)	d)
Compound mixture	5 %	25 %	50 %	80 %
Sodium ligninsulfonate	4 %	-	3 %	-
Sodium lauryl sulfate	2 %	3 %	-	4 %

- 18 -

Sodium diisobutylnaphthalene sulfonate -	6 %	5 %	6 %
Octylphenol polyethoxylate	1 %	2 %	-
(7-8 mols EO)			
Highly dispersed silicic acid	1 %	3 %	5 %
Kaolin	88 %	62 %	35 %

The compound is thoroughly mixed with the adjuvants and this mixture is ground in a suitable mill to give wettable powders that can be diluted with water to give suspensions of any desired concentration.

<u>F4. Coated granulates</u>	a)	b)	c)
Compound mixture	0.1 %	5 %	15 %
Highly dispersed silicic acid	0.9 %	2 %	2 %
Inorganic carrier	99.0 %	93 %	83 %

(Ø 0.1–1 mm)

such as CaCO_3 or SiO_2

The compound mixture is dissolved in dichloromethane, the solution is sprayed on to the carrier, and the solvent is removed under vacuum.

<u>F5. Coated granulates</u>	a)	b)	c)
Compound mixture	0.1 %	5 %	15 %
Polyethylene glycol mw 200	1.0 %	2 %	3 %
Highly dispersed silicic acid	0.9 %	1 %	2 %
Inorganic carrier	98.0 %	92 %	80 %

(Ø 0.1–1 mm)

such as CaCO_3 or SiO_2

The finely ground active substance is uniformly applied in a mixer to the carrier moistened with polyethylene glycol. Non-dusty coated granulates are obtained in this manner.

<u>F6. Extruder granulates</u>	a)	b)	c)	d)
Compound mixture	0.1 %	3 %	5 %	15 %
Sodium ligninsulfonate	1.5 %	2 %	3 %	4 %
Carboxymethylcellulose	1.4 %	2 %	2 %	2 %
Kaolin	97.0 %	93 %	90 %	79 %

- 19 -

The compound is mixed and ground with the adjuvants, and the mixture is moistened with water. This mixture is extruded and then dried in a stream of air.

<u>F7. Dusts</u>	a)	b)	c)
Compound mixture	0.1 %	1 %	5 %
Talcum or champagne chalk	39.9%	49 %	35 %
Kaolin	60.0 %	50 %	60 %

Ready-to-use dusts are obtained by mixing the active ingredient with the carriers and grinding on a suitable mill.

<u>F8. Suspension concentrates</u>	a)	b)	c)	d)
Compound mixture	3 %	10 %	25 %	50 %
Ethylene glycol	5 %	5 %	5 %	5 %
Nonylphenol polyethoxylate (15 mols EO)	-	1 %	2 %	-
Sodium ligninsulfonate	3 %	3 %	4 %	5 %
Carboxymethylcellulose	1 %	1 %	1 %	1 %
37% aqueous formaldehyde solution	0.2 %	0.2 %	0.2 %	0.2 %
Silicone oil emulsion	0.8 %	0.8 %	0.8 %	0.8 %
Water	87 %	79 %	62 %	38 %

The finely ground active substance is intimately mixed with the adjuvants. In this way, a suspension concentrate is obtained from which suspensions of any desired concentration can be prepared by dilution with water.

It is often more convenient to formulate the active ingredient of formula I and the compound(s) 2.1 to 2.33 separately and not to combine them until shortly before application in the applicator in the desired mixing ratio in the form of a "tank mix" in water.

Biological Examples:

A synergistic effect is always obtained when the herbicidal action of the combination of compound I and 2.1 to 2.33 is greater than the sum of the action of the individual herbicides.

- 20 -

The expected herbicidal action We for a given combination of two herbicides can be calculated as follows (q.v. COLBY, S.R., "Calculating synergistic and antagonistic response of herbicide combinations", Weeds 15, pages 20-22, 1967):

$$We = X + [Y \bullet (100 - X) / 100]$$

In this formula:

X = the percentage of herbicidal action after treatment with the compound of formula I at a rate of application of p kg per hectare, compared with untreated controls ($= 0\%$).

Y = the percentage of herbicidal action after treatment with a compound of formula 2.1 to 2.33 at a rate of application of q kg per hectare, compared with untreated controls.

We = the expected herbicidal action (percentage of herbicidal action compared with the untreated control plants) after treatment with the compounds of formulae I and 2.1 to 2.33 at a rate of application of $p + q$ kg per hectare.

If the value actually observed is higher than the expected value We , then there is synergism.

The synergistic effect of the combinations of the compound of formula I with the compounds of formulae 2.1 to 2.33 is demonstrated in the following Examples.

Description of postemergence test

The test plants are grown under greenhouse conditions in plastic pots until reaching the 2-3 leaf stage. Standard soil is used as the substrate for cultivation. At the 2-3 leaf stage, the herbicides are applied to the test plants on their own and in a mixture. Application is effected in the form of an aqueous suspension of the test substances in 500 l water/ha. The rates of application depend on the optimum dosages determined under field conditions and greenhouse conditions. Evaluation of the tests is made after 20 days (% activity, 100% = plant perished, 0% = no phytotoxic effect). The results are given in Tables 1 to 5 which follow.

The compound of formula I, wherein A₁ is nitro and A₂ is hydrogen, is designated as a compound of formula Ia. The compound of formula I, wherein A₁ is methyl and A₂ is methoxy, is designated as a compound of formula Ib.

Table 1: Synergistic effect of the mixture of the compound of formula Ia with halosulfuron:

Test plant	Ia 75 g/ha	halosulfuron 100 g/ha	Ia 75 g/ha + halosulfuron 100 g/ha	expected activity We according to Colby
maize DK 261	0	5	0	5
Digitaria	50	25	70	63

Table 2: Synergistic effect of the mixture of the compound of formula Ia with rimsulfuron:

Test plant	Ia 150 g/ha	rimsulfuron 30 g/ha	Ia 150 g/ha + rimsulfuron 30 g/ha	expected activity We according to Colby
maize DK 261	5	10	5	14,5
Rottboellia	50	25	70	63

Table 3: Synergistic effect of the mixture of the compound of formula Ia with glyphosate:

Test plant	Ia 150 g/ha	glyphosate 500 g/ha	Ia 150 g/ha + glyphosate 500 g/ha	expected activity We according to Colby
Cyperus	60	75	95	90

Table 4: Synergistic effect of the mixture of the compound of formula Ia with primisulfuron:

Test plant	Ia 75 g/ha	primisulfuron 20 g/ha	Ia 75 g/ha + primisulfuron 20 g/ha	expected activity We according to Colby
Echinochloa	80	10	85	82

Table 5: Synergistic effect of the mixture of the compound of formula Ib with glyphosate:

Test plant	Ib 150 g/ha	glyphosate 500 g/ha	Ib 150 g/ha + glyphosate 500 g/ha	expected activity We according to Colby
Cyperus	60	75	95	90

Description of pre-emergence test

The test plants are sown in plastic pots in standard soil under greenhouse conditions.

Directly after sowing, the test substances are applied in an aqueous suspension

(500 l water/ha). The test plants are then further cultivated in the greenhouse under optimum conditions. The rates of application depend on the optimum dosages determined under field conditions and greenhouse conditions. Evaluation of the tests is made after 26 days (% activity, 100% = plant perished, 0% = no phytotoxic effect). The results are given in Tables 6 to 9 which follow.

Table 6: Synergistic effect of the mixture of the compound of formula Ia with terbutylazin:

Test plant	Ia 100 g/ha	terbutylazin 200 g/ha	Ia 100 g/ha + terbutylazin 200 g/ha	expected activity We according to Colby
Ipomoea	75	30	95	82.5
Polygonum	60	60	100	84
Xanthium	80	0	95	80

Table 7: Synergistic effect of the mixture of the compound of formula Ia with atrazine:

Test plant	Ia 50 g/ha	atrazine 200 g/ha	Ia 50 g/ha + atrazine 200 g/ha	expected activity We according to Colby
Convolvulus	75	80	100	95
Polygonum	40	90	100	94
Xanthium	60	0	98	60

Table 8: Synergistic effect of the mixture of the compound of formula Ib with terbutylazin:

Test plant	Ib 50 g/ha	terbutylazin 200 g/ha	Ib 50 g/ha + terbutylazin 200 g/ha	expected activity We according to Colby
Convolvulus	60	80	98	92
Ipomoea	60	30	75	72
Polygonum	30	60	100	72
Xanthium	70	0	98	70

Table 9: Synergistic effect of the mixture of the compound of formula Ib with atrazine:

Test plant	Ib 50 g/ha	atrazine 200 g/ha	Ib 50 g/ha + atrazine 200 g/ha	expected activity We according to Colby
Convolvulus	60	80	98	92
Ipomoea	60	70	100	88
Polygonum	30	90	100	93
Xanthium	70	0	100	70

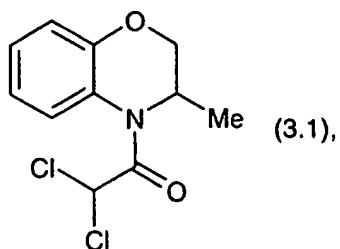
It has surprisingly been found that special safeners are suitable for mixing with the synergistic composition according to the invention. The present invention therefore also relates to a selective herbicidal composition to control grasses and weeds in crops of cultivated plants, especially maize, which contains a compound of formula I, one or more compounds selected from the compounds of formulae 2.1 to 2.33 and a safener (antidote) and to protect the cultivated plants, but not the weeds, from the phytotoxic action of the herbicide, and to the use of said composition for controlling weeds in crops of cultivated plants

Accordingly, the invention also provides a selective herbicidal composition comprising, in addition to customary inert formulation assistants such as carriers, solvents and wetting agents, a mixture of

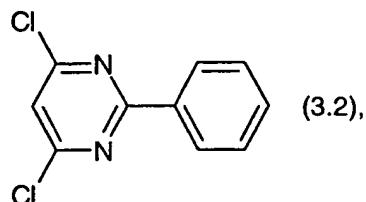
a) herbicidally synergistic amount of a compound of formula I and one or more compounds selected from the compounds of formulae 2.1 to 2.33 and

- 25 -

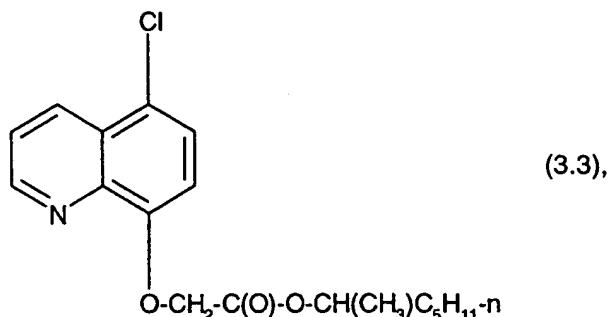
b) to antagonise the herbicide, an antidotally effective amount of a safener selected from the compound of formula 3.1



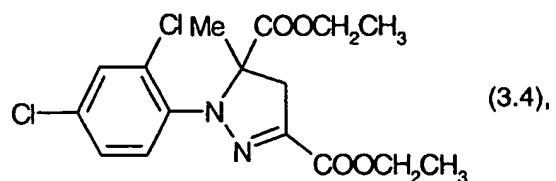
and the compound of formula 3.2



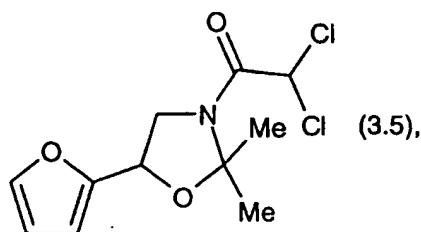
and the compound of formula 3.3



and the compound of formula 3.4

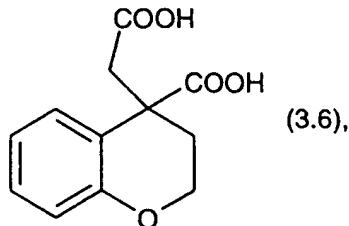


and the compound of formula 3.5

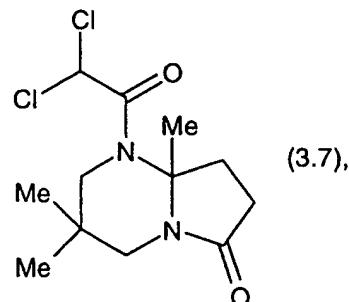


- 26 -

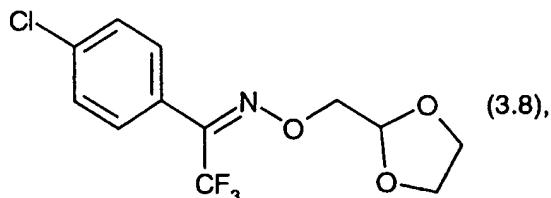
and the compound of formula 3.6



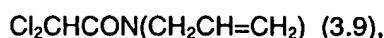
and the compound of formula 3.7



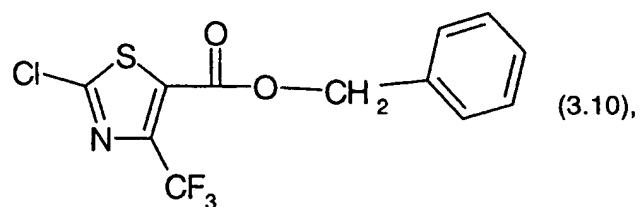
and the compound of formula 3.8



and of formula 3.9

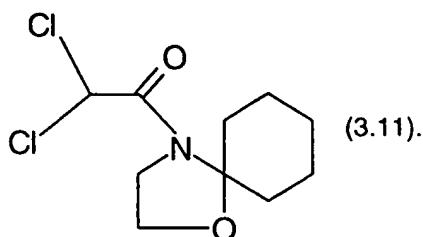


and of formula 3.10

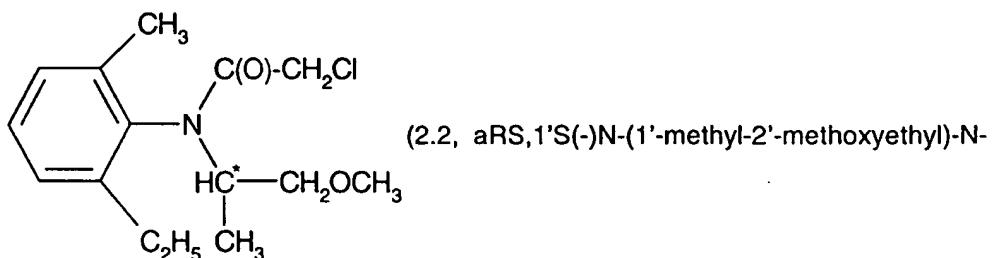


and of formula 3.11

- 27 -

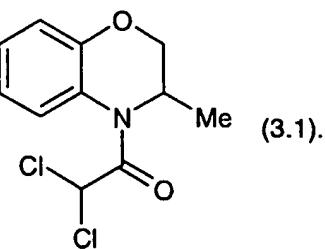


Preferred selective herbicidal compositions contain a) a herbicidally synergistic amount of a compound of formula I, a compound of formula 2.2

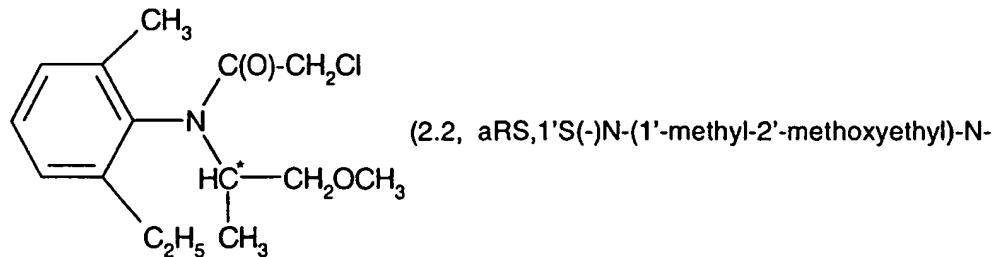


chloroacetyl-2-ethyl-6-methylaniline), as well as a compound selected from formula 2.4, wherein R₇ is chlorine, R₈ is ethyl and R₉ is isopropyl, or R₇ is chlorine, R₈ is ethyl and R₉ is tert.-butyl, and of formula 2.6, wherein R₁₂ is hydrogen, Z is methine, R₁₃ is methyl, Y is nitrogen, R₁₄ is fluorine, R₁₅ is hydrogen and R₁₆ is fluorine, or R₁₂ is methoxy, Z is methine, R₁₃ is methoxy, Y is methine, R₁₄ is chlorine, R₁₅ is methyl and R₁₆ is chlorine, and of formula 2.7, wherein R₁₇ is -C(O)-S-n-octyl, and of formula 2.13, wherein Y₁ is nitrogen, Y₂, Y₃ and Y₄ are methine, R₂₄ is dimethylaminocarbonyl and Y₅ is methine, or Y₁, Y₂, Y₃ and Y₄ are methine, R₂₄ is methoxycarbonyl and Y₅ is methine, or Y₁, Y₂, Y₃ and Y₄ are methine, R₂₄ is 3-trifluoropropyl and Y₅ is nitrogen, or Y₁ is nitrogen, Y₂, Y₃ and Y₄ are methine, R₂₄ is ethylsulphonyl and Y₅ is methine, or Y₁ is N-Me, Y₂ is nitrogen, Y₃ and Y₄ together are C-Cl, R₂₄ is methoxycarbonyl and Y₅ is methine, and of formula 2.9, and of formula 2.11, wherein R₂₂ is trifluoromethyl, and of formula 2.12, and of formula 2.16, and of formula 2.18, and of formula 2.19, and

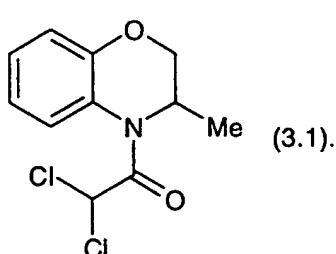
b) to antagonise the herbicide, an antidotally effective amount of a safener of formula 3.1



An especially preferred selective herbicidal composition contains a) a herbicidally synergistic amount of a compound of formula I and a compound of formula 2.2

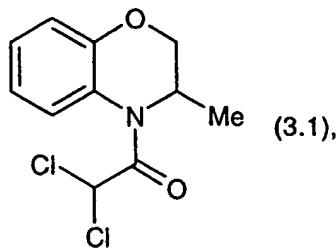


b) to antagonise the herbicide, an antidotally effective amount of a safener of formula 3.1

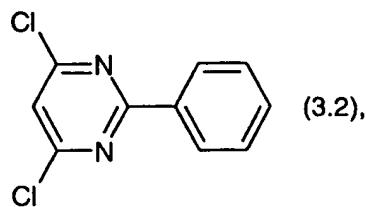


The invention also provides a selective herbicidal composition comprising, in addition to customary inert formulation assistants such as carriers, solvents and wetting agents, a mixture of

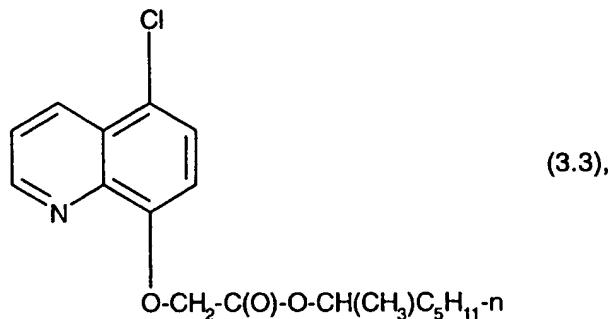
a) a herbicidally effective amount of a compound of formula I and
 b) to antagonise the herbicide, an antidotally effective amount of a safener selected from the compound of formula 3.1



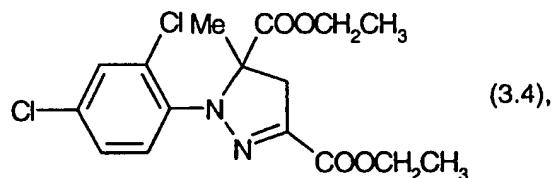
and the compound of formula 3.2



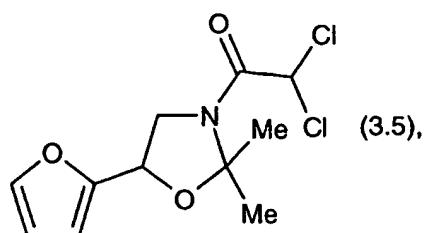
and the compound of formula 3.3



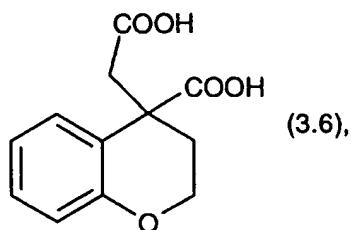
and the compound of formula 3.4



and the compound of formula 3.5

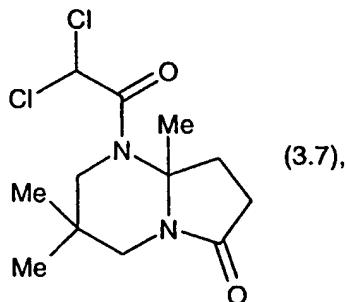


and the compound of formula 3.6

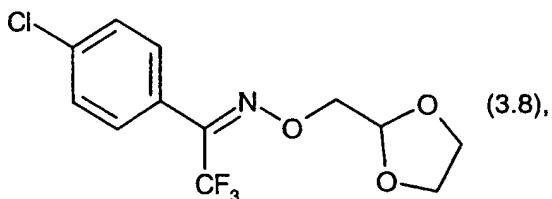


and the compound of formula 3.7

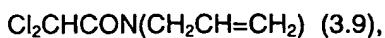
- 30 -



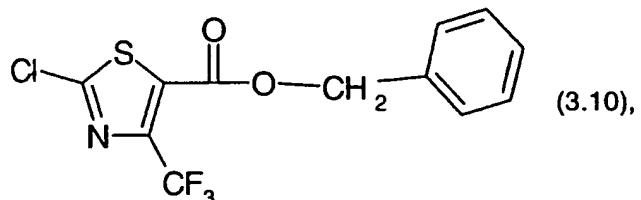
and the compound of formula 3.8



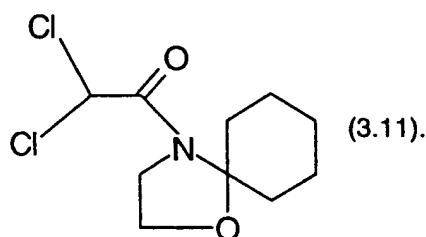
and of formula 3.9



and of formula 3.10



and of formula 3.11



The invention additionally relates to a process for the selective control of weeds in crops of cultivated plants, which comprises treating said cultivated plants, the seeds or seedlings or the crop area thereof with a herbicidally effective amount of the herbicide of formula I, if required, one or more herbicides selected from the compounds of formulae 2.1 to 2.33 and,

to antagonise the herbicide, an antidotally effective amount of a safener of formula 3.1 to 3.11.

The compounds of formulae 3.1 to 3.11 are known and are described for example in the Pesticide Manual, eleventh ed., British Crop Protection Council, 1997 under the entry nos. 61 (formula 3.1, benoxacor), 304 (formula 3.2, fenclorim), 154 (formula 3.3, cloquintocet), 462 (formula 3.4, mefenpyr-diethyl), 377 (formula 3.5, furilazol), 363 (formula 3.8, fluxofenim), 213 (formula 3.9, dichlormid) and 350 (formula 3.10, flurazole). The compound of formula 3.11 is known under the designation MON 4660 (Monsanto).

The compound of formula 3.6 (AC 304415) is described for example in EP-A-0 613.618, and the compound of formula 3.7 is described in DE-A-2948535.

Crop plants which may be protected against the harmful action of the above-mentioned herbicides by the safeners of formulae 3.1 to 3.11 are in particular cereals, cotton, soya, sugar beet, sugar cane, plantations, rape, maize and rice, especially maize. Crops will also be understood as meaning those crops that have been made tolerant to herbicides or classes of herbicides by conventional breeding or genetic engineering methods.

The weeds to be controlled may be monocot as well as dicot weeds, typically *Stellaria*, *Agrostis*, *Digitaria*, *Avena*, *Apera*, *Brachiaria*, *Phalaris*, *Setaria*, *Sinapis*, *Lolium*, *Solanum*, *Echinochloa*, *Scirpus*, *Monochoria*, *Sagittaria*, *Panicum*, *Bromus*, *Alopecurus*, *Sorghum halepense*, *Sorghum bicolor*, *Rottboellia*, *Cyperus*, *Abutilon*, *Sida*, *Xanthium*, *Amaranthus*, *Chenopodium*, *Ipomoea*, *Chrysanthemum*, *Galium*, *Viola*, and *Veronica*.

Crop areas will be understood as meaning the areas already under cultivation with the cultivated plants or seeds thereof, as well as the areas intended for cropping with said cultivated plants.

Depending on the end use, a safener of formula 3.1 to 3.11 can be used for pretreating seeds of the crop plants (dressing of seeds or seedlings) or it can be incorporated in the soil before or after sowing. It can, however, also be applied by itself alone or together with the herbicide postemergence. Treatment of the plant or the seeds with the safener can therefore in principle be carried out irrespective of the time of application of the herbicide.

Treatment can, however, also be carried out by simultaneous application of the herbicide and safener (e.g. as tank mixture). The concentration of safener with respect to the herbicide will depend substantially on the mode of application. Where a field treatment is carried out either by using a tank mixture with a combination of safener and herbicide or by separate application of safener and herbicide, the ratio of herbicide to safener will usually be from 100:1 to 1:10, preferably 20:1 to 1:1. In field treatment it is usual to apply 0.001 to 1.0 kg/ha, preferably 0.001 to 0.25 kg/ha, of safener.

The concentration of herbicide is usually in the range from 0.001 to 2 kg/ha, but will preferably be from 0.005 to 0.5 kg/ha.

The compositions of this invention are suitable for all methods of application commonly used in agriculture, including preemergence application, postemergence application and seed dressing.

For seed dressing, 0.001 to 10 g of safener/kg of seeds, preferably 0.05 to 2 g of safener/kg of seeds, is usually applied. If the safener is used in liquid form shortly before sowing to effect soaking, then it is preferred to use safener solutions that contain the active ingredient in a concentration of 1 to 10000 ppm, preferably of 100 to 1000 ppm.

For application, it is preferred to process the safeners of formula 3.1 to 3.11, or mixtures of these safeners with the herbicide of formula I and optionally with one or more herbicides selected from formulae 2.1 to 2.33, conveniently together with the customary assistants of formulation technology to formulations, typically to emulsifiable concentrates, coatable pastes, directly sprayable or dilutable solutions, dilute emulsions, wettable powders, soluble powders, dusts, granulates or microcapsules.

Such formulations are described, for example, in WO 97/34485 on pages 9 to 13. The formulations are prepared in known manner, conveniently by homogeneously mixing and/or grinding the active ingredients with liquid or solid formulation assistants, typically solvents or solid carriers. Surface-active compounds (surfactants) may additionally be used for preparing the formulations. Solvents and solid carriers that are suitable for this purpose are described in WO 97/34485 on page 6.

Depending on the herbicide of formula I, 2.1 to 2.33 and 3.1 to 3.11 to be formulated, suitable surface-active compounds are nonionic, cationic and/or anionic surfactants and surfactant mixtures having good emulsifying, dispersing and wetting properties. Examples of suitable anionic, nonionic, and cationic surfactants are listed in WO 97/34485 on pages 7 and 8. Also the surfactants customarily used for the art of formulation and described, *inter alia*, in "Mc Cutcheon's Detergents and Emulsifiers Annual" MC Publishing Corp., Ridgewood New Jersey, 1981, Stache, H., "Tensid-Taschenbuch" (Handbook of Surfactants), Carl Hanser Verlag, Munich/Vienna, 1981, and M. and J. Ash, "Encyclopedia of Surfactants", Vol I-III, Chemical Publishing Co., New York, 1980-81 are suitable for manufacture of the herbicides according to the invention.

The herbicidal compositions will usually contain from 0.1 to 99% by weight, preferably from 0.1 to 95% by weight, of compound mixture of the compound of formula I, a compound selected from the compounds of formulae 2.1 to 2.33 and the compounds of formulae 3.1 to 3.11, from 1 to 99.9% by weight of a solid or liquid formulation assistant, and from 0 to 25% by weight, preferably from 0.1 to 25% by weight, of a surfactant. Whereas it is customarily preferred to formulate commercial products as concentrates, the end user will normally use dilute formulations.

The compositions may also contain further ingredients, such as stabilisers, e.g. where appropriate epoxidised vegetable oils (epoxidised coconut oil, rapeseed oil, or soybean oil), antifoams, typically silicone oil, preservatives, viscosity regulators, binders, tackifiers, as well as fertilisers or other chemical agents. Different methods and techniques may suitably be used for applying the safeners of formula 3.1 to 3.11 or compositions containing them to protect cultivated plants from the harmful effects of herbicides of formula I and 2.1 to 2.33, for example the following:

- i) Seed dressing
 - a) Dressing the seeds with a wettable powder formulation of the active ingredient of formulae 3.1 to 3.11 by shaking in a vessel until the safener is uniformly distributed on the surface of the seeds (dry treatment). In this instance, approximately 1 to 500 g of active ingredient of formula 3.1 to 3.11 (4 g to 2 kg of wettable powder) is used per 100 kg of seeds.

- b) Dressing seeds with an emulsifiable concentrate of the active ingredient of formulae 3.1 to 3.11 by method a) (wet treatment).
- c) Dressing by immersing the seeds in a mixture containing 100–1000 ppm of active ingredient of formulae 3.1 to 3.11 for 1 to 72 hours and where appropriate subsequently drying them (seed soaking).

In keeping with the natural environment, the preferred method of application is either seed dressing or treatment of the germinated seedlings, because the safener treatment is fully concentrated on the target crop. Usually 1 to 1000 g, preferably 5 to 250 g, of safener is used per 100 kg of seeds. However, depending on the method employed, which also permits the use of other chemical agents or micronutrients, the concentrations may deviate above or below the indicated limit values (repeat dressing).

ii) Application as a tank mixture

A liquid formulation of a mixture of safener and herbicide (reciprocal ratio from 10:1 to 1:100) is used, the concentration of herbicide being from 0.005 to 5.0 kg/ha. This tank mixture is applied before or after sowing.

iii) Application in the furrow

The active ingredients of formulae 3.1 to 3.11 formulated as emulsifiable concentrate, wettable powder or granulate are applied to the open furrow in which the seeds have been sown. After covering the furrow, the herbicide is applied pre-emergence in conventional manner.

iv) Controlled release of compound

A solution of the active ingredients of formulae 3.1 to 3.11 is applied to a mineral granular carrier or to a polymerised granulate (urea/formaldehyde) and then dried. A coating can then be applied (coated granules) that allows the herbicide to be released at a controlled rate over a specific period of time.

Particularly preferred formulations are made up as follows:

(% = percent by weight)

Emulsifiable concentrates:

Compound mixture: 1 to 90 %, preferably 5 to 20 %

Surfactant: 1 to 30 %, preferably 10 to 20 %

Liquid carrier: 5 to 94 %, preferably 70 to 85 %

Dusts:

Compound mixture: 0.1 to 10 %, preferably 0.1 to 5 %

Solid carrier: 99.9 to 90 %, preferably 99.9 to 99 %

Suspension concentrates:

Compound mixture: 5 to 75 %, preferably 10 to 50 %

Water: 94 to 24 %, preferably 88 to 30 %

Surfactant: 1 to 40 %, preferably 2 to 30 %

Wettable powders:

Compound mixture: 0.5 to 90 %, preferably 1 to 80 %

Surfactant: 0.5 to 20 %, preferably 1 to 15 %

Solid carrier: 5 to 95 %, preferably 15 to 90 %

Granulates:

Compound mixture: 0.1 to 30 %, preferably 0.1 to 15 %

Solid carrier: 99.5 to 70 %, preferably 97 to 85 %

The invention is illustrated by the following non-limitative Examples.

Formulation examples for mixtures of herbicides of formula I, formulae 2.1 to 2.33 and safeners of formulae 3.1 to 3.11 (% = percent by weight)

<u>F1. Emulsifiable concentrates</u>	a)	b)	c)	d)
Compound mixture	5 %	10 %	25 %	50 %
Calcium dodecylbenzenesulfonate	6 %	8 %	6 %	8 %
Polyethoxylated castor oil (36 mols EO)	4 %	-	4 %	4 %
Octylphenol polyethoxylate (7-8 mols EO)	-	4 %	-	2 %
Cyclohexanone	-	-	10 %	20 %

- 36 -

Aromatic hydrocarbon mixture C ₉ -C ₁₂	85 %	78 %	55 %	16 %
--	------	------	------	------

Emulsions of any desired concentration can be prepared by diluting such concentrates with water.

<u>F2. Solutions</u>	a)	b)	c)	d)
Compound mixture	5 %	10 %	50 %	90 %
1-methoxy-3-(3-methoxy-propoxy)-propane	-	20 %	20 %	-
Polyethylene glycol mw 400	20 %	10 %	-	-
N-Methyl-2-pyrrolidone	-	-	30 %	10 %
Aromatic hydrocarbon mixture C ₉ -C ₁₂	75 %	60 %	-	-

The solutions are suitable for use in the form of microdrops.

<u>F3. Wettable powders</u>	a)	b)	c)	d)
Compound mixture	5 %	25 %	50 %	80 %
Sodium ligninsulfonate	4 %	-	3 %	-
Sodium lauryl sulfate	2 %	3 %	-	4 %
Sodium diisobutylnaphthalene sulfonate	-	6 %	5 %	6 %
Octylphenol polyethoxylate (7-8 mols EO)	-	1 %	2 %	-
Highly dispersed silicic acid	1 %	3 %	5 %	10 %
Kaolin	88 %	62 %	35 %	-

The compound is thoroughly mixed with the adjuvants and this mixture is ground in a suitable mill to give wettable powders which can be diluted with water to give suspensions of any desired concentration.

<u>F4. Coated granulates</u>	a)	b)	c)
Compound mixture	0.1 %	5 %	15 %
Highly dispersed silicic acid	0.9 %	2 %	2 %
Inorganic carrier (Ø 0.1-1 mm)	99.0 %	93 %	83 %

such as CaCO_3 or SiO_2

The compound mixture is dissolved in dichloromethane, the solution is sprayed on to the carrier, and the solvent is removed under vacuum.

- 38 -

<u>F5. Coated granulates</u>	a)	b)	c)
Compound mixture	0.1 %	5 %	15 %
Polyethylene glycol mw 200	1.0 %	2 %	3 %
Highly dispersed silicic acid	0.9 %	1 %	2 %
Inorganic carrier	98.0 %	92 %	80 %

(Ø 0.1–1 mm)

such as CaCO₃ or SiO₂

The finely ground active substance is uniformly applied in a mixer to the carrier moistened with polyethylene glycol. Non-dusty coated granulates are obtained in this manner.

<u>F6. Extruder granulates</u>	a)	b)	c)	d)
Compound mixture	0.1 %	3 %	5 %	15 %
Sodium ligninsulfonate	1.5 %	2 %	3 %	4 %
Carboxymethylcellulose	1.4 %	2 %	2 %	2 %
Kaolin	97.0 %	93 %	90 %	79 %

The compound is mixed and ground with the adjuvants, and the mixture is moistened with water. This mixture is extruded and then dried in a stream of air.

<u>F7. Dusts</u>	a)	b)	c)
Compound mixture	0.1 %	1 %	5 %
Talc	39.9 %	49 %	35 %
Kaolin	60.0 %	50 %	60 %

Ready-to-use dusts are obtained by mixing the active ingredient with the carriers and grinding on a suitable mill.

<u>F8. Suspension concentrates</u>	a)	b)	c)	d)
Compound mixture	3 %	10 %	25 %	50 %
Ethylene glycol	5 %	5 %	5 %	5 %
Nonylphenol polyethoxylate (15 mols EO)	-	1 %	2 %	-
Sodium ligninsulfonate	3 %	3 %	4 %	5 %
Carboxymethylcellulose	1 %	1 %	1 %	1 %
37% aqueous formaldehyde	0.2 %	0.2 %	0.2 %	0.2 %

solution

Silicone oil emulsion	0.8 %	0.8 %	0.8 %	0.8 %
Water	87 %	79 %	62 %	38 %

The finely ground active substance is intimately mixed with the adjuvants. In this way, a suspension concentrate is obtained, from which suspensions of any desired concentration can be prepared by dilution with water.

It is often more convenient to formulate the active ingredients of formulae I, 2.1 to 2.33 and 3.1 to 3.11 separately and not to combine them until shortly before application in the applicator in the desired mixing ratio in the form of a "tank mix" in water.

The ability of the safeners of formulae 3.1 to 3.11 to protect crops from the phytotoxic action of herbicides of formula I will be demonstrated in the examples which follow.

Biological Example: Safening effect

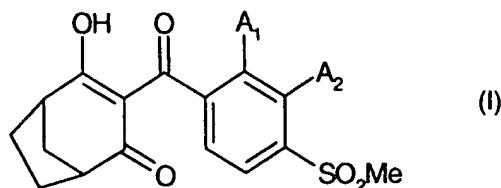
The test plants are grown under greenhouse conditions in plastic pots until reaching the 4 leaf stage. At this stage, the herbicides are applied to the test plants both on their own and in mixtures with the test substances that are to be tested as safeners. Application is effected in the form of an aqueous suspension of the test substances, prepared from a 25% wettable powder [example F3, b)] at 500 l water/ha. 3 weeks after application, the phytotoxic effect of the herbicides on the cultivated plants, such as maize and cereals, is evaluated on a percentage scale. 100% indicates that the test plant has perished, 0% indicates no phytotoxic effect.

The results obtained in this test show that the damage to the cultivated plant caused by the herbicide of formula I in combination with one or more herbicides selected from formulae 2.1 to 2.33 can be significantly reduced with the compounds of formulae 3.1 to 3.11.

The same results are obtained by formulating the mixtures in accordance with Examples F1, F2 and F4 to F8.

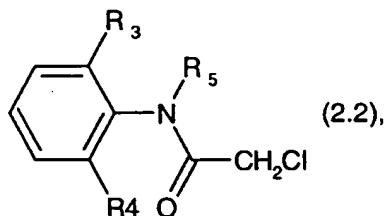
What is claimed is:

1. A herbicidal synergistic composition comprising, in addition to customary inert formulation assistants, as the active ingredient, a combination of
 a) a compound of formula I



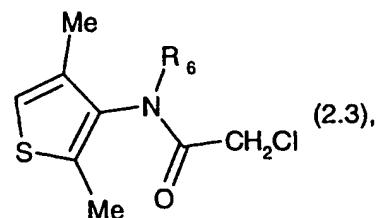
wherein either A₁ is nitro and A₂ is hydrogen or A₁ is methyl and A₂ is methoxy, as well as their salts, and

b) a synergistically active amount of one or more compounds selected from the compound of formula 2.2



wherein R₃ is ethyl, R₄ is methyl or ethyl and R₅ is -CH(Me)-CH₂OMe, <S>-CH(Me)-CH₂OMe, CH₂OMe or CH₂O-CH₂CH₃;

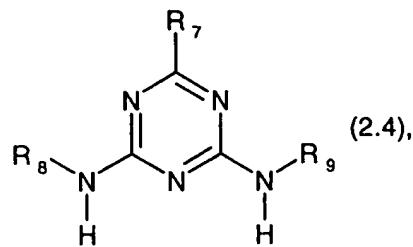
and the compound of formula 2.3



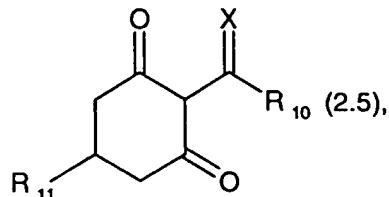
wherein R₆ is CH(Me)-CH₂OMe or <S>CH(Me)-CH₂OMe;

and the compound of formula 2.4

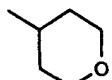
- 41 -



wherein R₇ is chlorine or SMe, R₈ is ethyl and R₉ is ethyl, isopropyl or tert.-butyl;
and the compound of formula 2.5

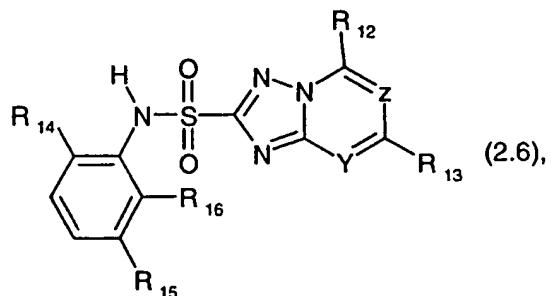


wherein R₁₀ is ethyl or n-propyl, R₁₁ is COO⁻ 1/2 Ca⁺⁺, -CH₂-CH(Me)S-CH₂CH₃ or the group



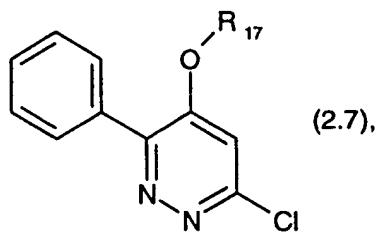
and X is oxygen, N-O-CH₂CH₃ or N-O-CH₂CH=CH-Cl;

and the compound of formula 2.6



wherein R₁₂ is hydrogen, methoxy or ethoxy, R₁₃ is Me, methoxy or fluorine, R₁₄ is COOMe, fluorine or chlorine, R₁₅ is hydrogen or Me, Y is methine or nitrogen, Z is methine or nitrogen and R₁₆ is fluorine or chlorine;

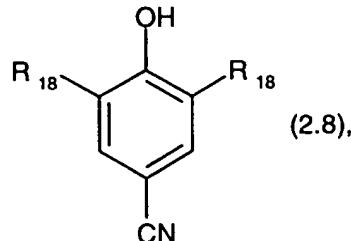
and the compound of formula 2.7



- 42 -

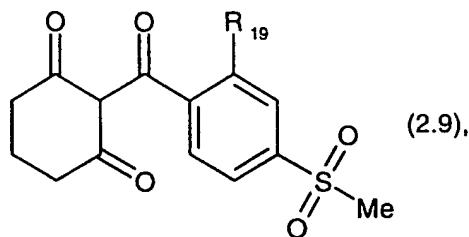
wherein R_{17} is hydrogen or $-C(O)-S-n\text{-octyl}$;

and the compound of formula 2.8



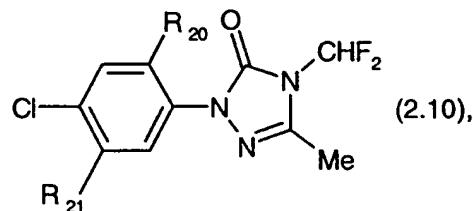
wherein R_{18} is bromine or iodine;

and the compound of formula 2.9



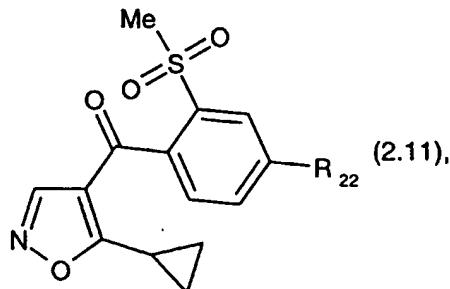
wherein R_{19} is chlorine or nitro;

and the compound of formula 2.10



wherein R_{20} is fluorine or chlorine and R_{21} is $-\text{CH}_2\text{CH}(\text{Cl})\text{COOCH}_2\text{CH}_3$ or $-\text{NH-SO}_2\text{Me}$;

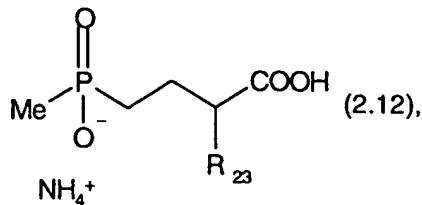
and the compound of formula 2.11



wherein R_{22} is trifluoromethyl or chlorine;

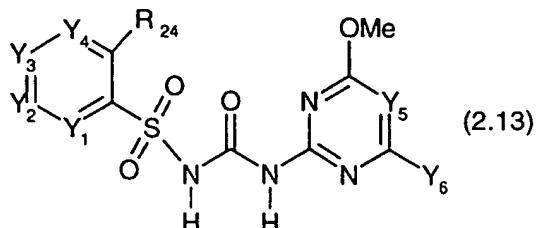
and the compound of formula 2.12

- 43 -



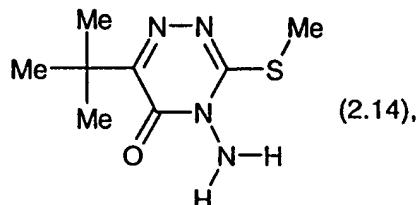
wherein R₂₃ is NH₂ or <S>NH₂;

and the compound of formula 2.13

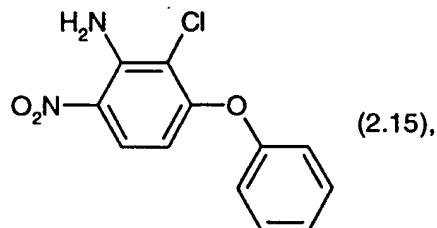


wherein Y₁ is nitrogen, methine or N-Me, Y₂ is nitrogen, methine or C-I, Y₃ is methine, Y₄ is methine, or Y₃ and Y₄ together are sulphur or C-Cl, Y₅ is nitrogen or methine, Y₆ is methyl or methoxy and R₂₄ is CONMe₂, COOMe, CH₂-CH₂CF₃ or SO₂CH₂CH₃, or the sodium salts thereof;

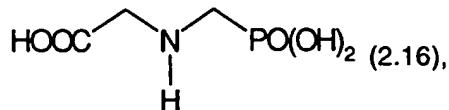
and the compound of formula 2.14



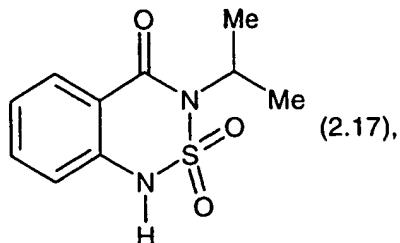
and the compound of formula 2.15



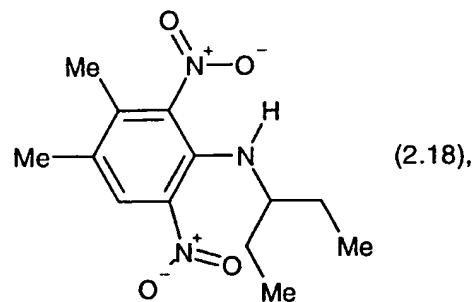
and the compound of formula 2.16



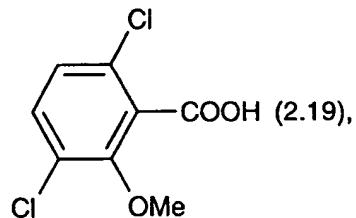
and the compound of formula 2.17



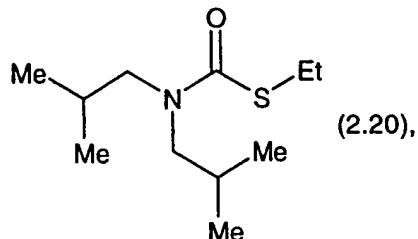
and the compound of formula 2.18



and the compound of formula 2.19

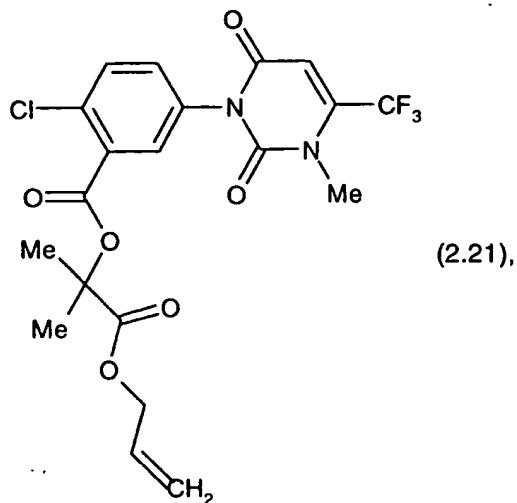


and the compound of formula 2.20

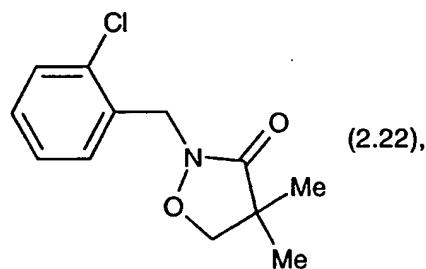


and the compound of formula 2.21

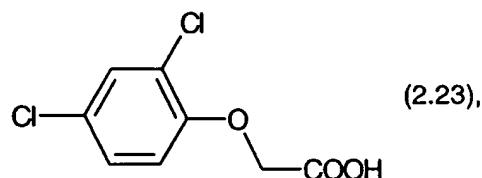
- 45 -



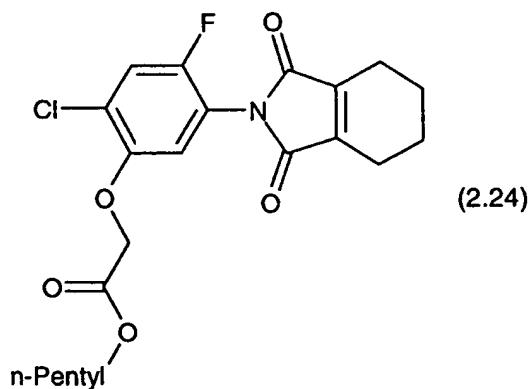
and the compound of formula 2.22



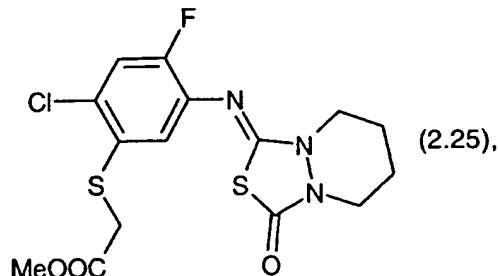
and the compound of formula 2.23



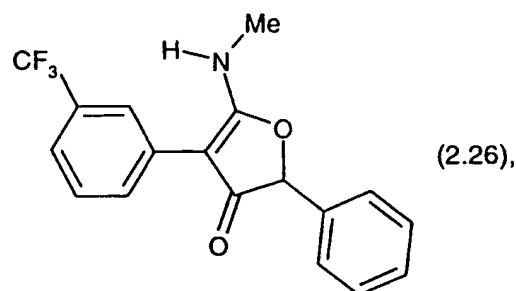
and the compound of formula 2.24



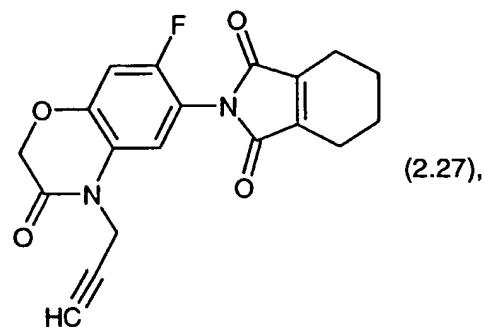
and the compound of formula 2.25



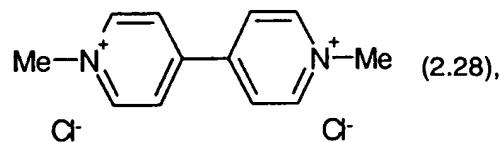
and the compound of formula 2.26



and the compound of formula 2.27

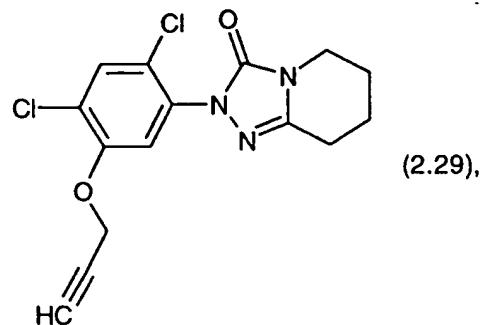


and the compound of formula 2.28

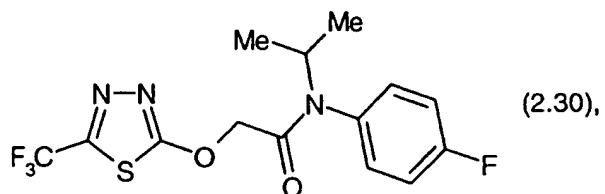


and the compound of formula 2.29

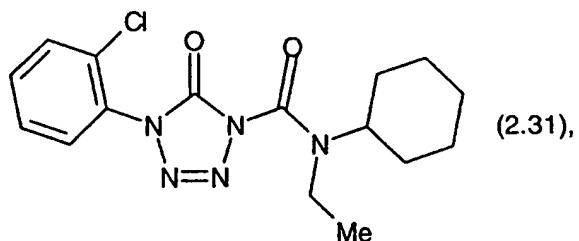
- 47 -



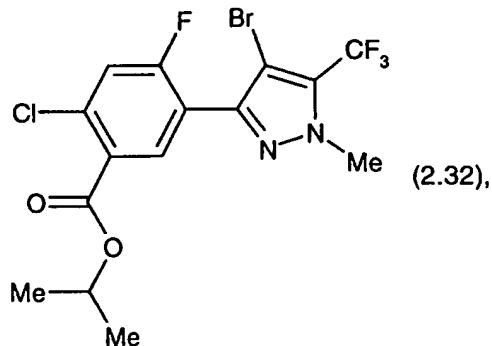
and the compound of formula 2.30



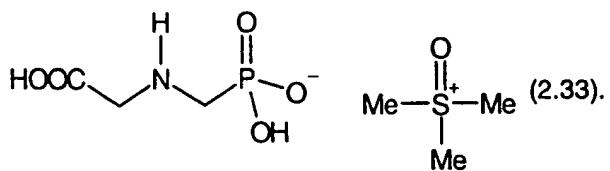
and the compound of formula 2.31



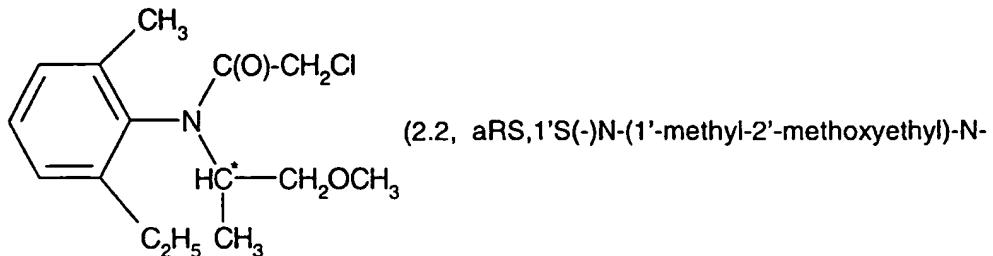
and the compound of formula 2.32



and the compound of formula 2.33

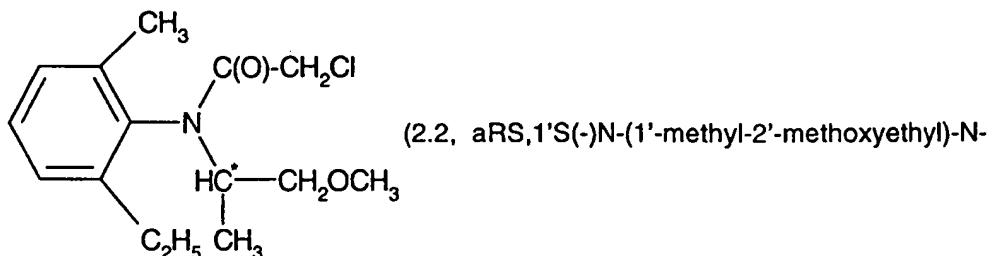


2. A herbicidal composition according to claim 1, comprising a compound of formula I and a synergistically effective amount either of a compound of formula 2.2



chloroacetyl-2-ethyl-6-methylaniline), or a compound of formula 2.3, or a compound of formula 2.4, wherein R₇ is chlorine, R₈ is ethyl and R₉ is isopropyl, or R₇ is chlorine, R₈ is ethyl and R₉ is tert.-butyl, or a compound of formula 2.6, wherein R₁₂ is hydrogen, Z is methine, R₁₃ is methyl, Y is nitrogen, R₁₄ is fluorine, R₁₅ is hydrogen and R₁₆ is fluorine, or R₁₂ is methoxy, Z is methine, R₁₃ is methoxy, Y is methine, R₁₄ is chlorine, R₁₅ is methyl and R₁₆ is chlorine, or a compound of formula 2.7, wherein R₁₇ is -C(O)-S-n-octyl, or a compound of formula 2.9, or a compound of formula 2.11, wherein R₂₂ is trifluoromethyl, or a compound of formula 2.12, or a compound of formula 2.13, wherein Y₁ is nitrogen, Y₂, Y₃ and Y₄ are methine, R₂₄ is dimethylaminocarbonyl and Y₅ is methine, or Y₁, Y₂, Y₃ and Y₄ are methine, R₂₄ is methoxycarbonyl and Y₅ is methine, or Y₁, Y₂, Y₃ and Y₄ are methine, R₂₄ is 3-trifluoropropyl and Y₅ is nitrogen, or Y₁ is nitrogen, Y₂, Y₃ and Y₄ are methine, R₂₄ is ethylsulphonyl and Y₅ is methine, or Y₁ is N-Me, Y₂ is nitrogen, Y₃ and Y₄ together are C-Cl, R₂₄ is methoxycarbonyl and Y₅ is methine, or a compound of formula 2.16, or a compound of formula 2.18, or a compound of formula 2.19, or a compound of formula 2.30.

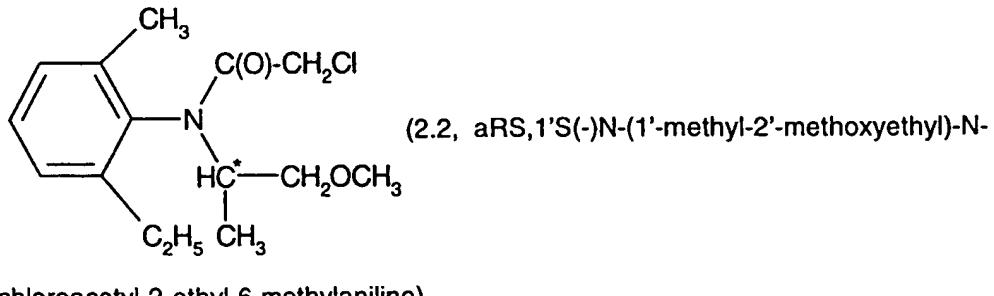
3. A herbicidal composition according to claim 1, comprising a compound of formula I, a compound of formula 2.2



chloroacetyl-2-ethyl-6-methylaniline), as well as a compound selected from formula 2.4, wherein R₇ is chlorine, R₈ is ethyl and R₉ is isopropyl, or R₇ is chlorine, R₈ is ethyl and R₉ is tert.-butyl, and of formula 2.6, wherein R₁₂ is hydrogen, Z is methine, R₁₃ is methyl, Y is

nitrogen, R₁₄ is fluorine, R₁₅ is hydrogen and R₁₆ is fluorine, or R₁₂ is methoxy, Z is methine, R₁₃ is methoxy, Y is methine, R₁₄ is chlorine, R₁₅ is methyl and R₁₆ is chlorine, and of formula 2.7, wherein R₁₇ is -C(O)-S-n-octyl, and of formula 2.13, wherein Y₁ is nitrogen, Y₂, Y₃ and Y₄ are methine, R₂₄ is dimethylaminocarbonyl and Y₅ is methine, or Y₁, Y₂, Y₃ and Y₄ are methine, R₂₄ is methoxycarbonyl and Y₅ is methine, or Y₁, Y₂, Y₃ and Y₄ are methine, R₂₄ is 3-trifluoropropyl and Y₅ is nitrogen, or Y₁ is nitrogen, Y₂, Y₃ and Y₄ are methine, R₂₄ is ethylsulphonyl and Y₅ is methine, or Y₁ is N-Me, Y₂ is nitrogen, Y₃ and Y₄ together are C-Cl, R₂₄ is methoxycarbonyl and Y₅ is methine, and of formula 2.9, and of formula 2.11, wherein R₂₂ is trifluoromethyl, and of formula 2.12, and of formula 2.16, and of formula 2.18, and of formula 2.19.

4. A herbicidal composition according to claim 1, comprising a compound of formula I and a synergistically effective amount of a compound of formula 2.2



5. A herbicidal composition according to claim 1, wherein the compound of formula I is present in a weight ratio to the compounds of formulae 2.1 to 2.33 of 1:2000 to 2000:1.

6. A method of controlling undesirable plant growth in crops of cultivated plants, which comprises treating said plants or the locus thereof with a herbicidally effective amount of a composition as claimed in claim 1.

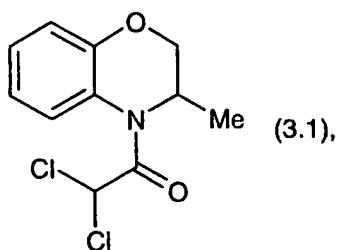
7. A method according to claim 6 wherein the cultivated plant is maize.

8. A method according to claim 6 which comprises treating the crops of cultivated plants with the said composition at rates of application corresponding to 1 to 5000 g total active ingredient per hectare.

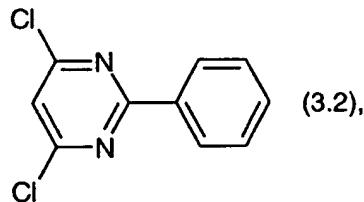
9. Selective herbicidal composition comprising, in addition to customary inert formulation assistants, such as carriers, solvents and wetting agents, as active ingredient a mixture of

a) herbicidally synergistic amount of a compound of formula 1 according to claim 1 and one or more compounds selected from the compounds of formulae 2.1 to 2.33 according to claim 1 and

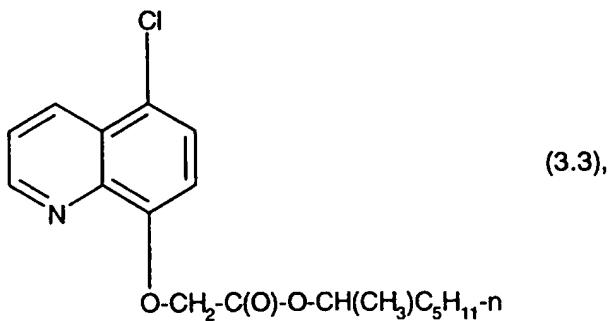
b) to antagonise the herbicide, an antidotally effective amount of a safener selected from the compound of formula 3.1



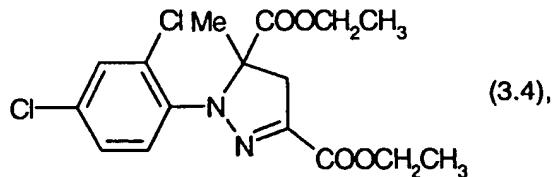
and the compound of formula 3.2



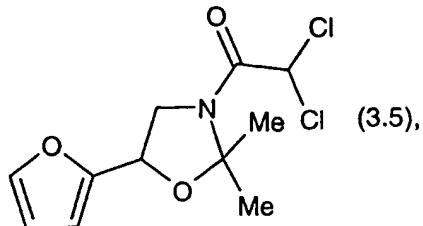
and the compound of formula 3.3



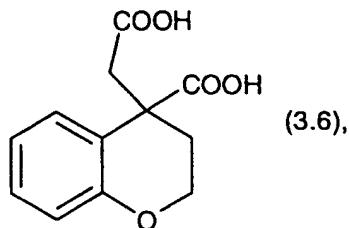
and the compound of formula 3.4



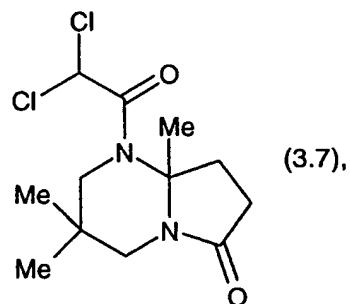
and the compound of formula 3.5



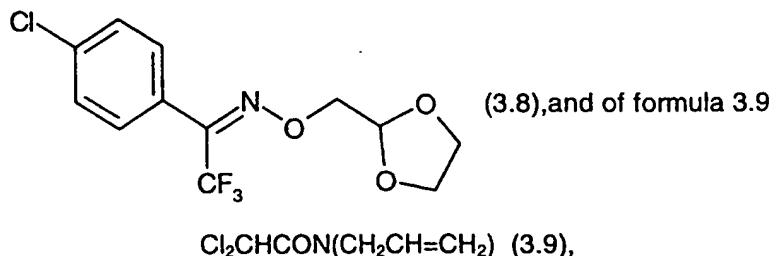
and the compound of formula 3.6



and the compound of formula 3.7

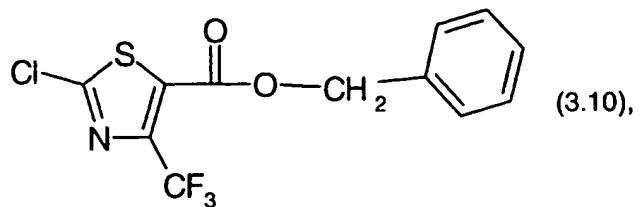


and the compound of formula 3.8

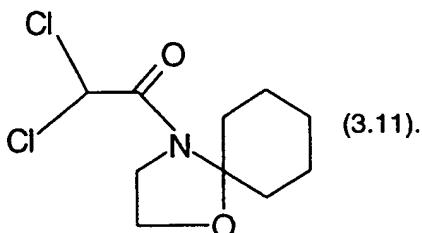


and of formula 3.10

- 52 -



and of formula 3.11

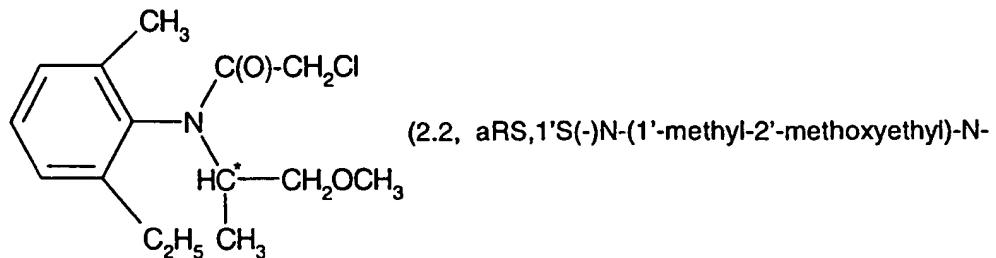


10. A method of selectively controlling weeds and grasses in crops of cultivated plants, which comprises treating said cultivated plants, the seeds or seedlings or the crop area thereof, with a herbicidally synergistic amount of a composition according to claim 9.

11. A method according to claim 10, wherein the rate of application of herbicides is 1 to 5000 g/ha and the rate of application of safeners is 0.001 to 0.5 kg/ha.

12. A method according to claim 10 wherein the cultivated plant is maize.

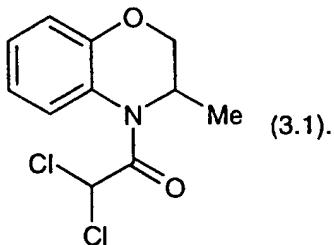
13. Herbicidal composition according to claim 9, which contains a) a herbicidally synergistic amount of a compound of formula I, a compound of formula 2.2



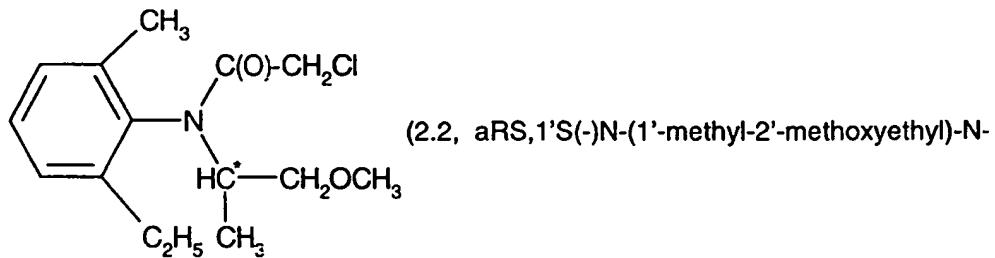
chloroacetyl-2-ethyl-6-methylaniline), as well as a compound selected from formula 2.4, wherein R₇ is chlorine, R₈ is ethyl and R₉ is isopropyl, or R₇ is chlorine, R₈ is ethyl and R₉ is tert.-butyl, and of formula 2.6, wherein R₁₂ is hydrogen, Z is methine, R₁₃ is methyl, Y is nitrogen, R₁₄ is fluorine, R₁₅ is hydrogen and R₁₆ is fluorine, or R₁₂ is methoxy, Z is methine, R₁₃ is methoxy, Y is methine, R₁₄ is chlorine, R₁₅ is methyl and R₁₆ is chlorine, and of

formula 2.7, wherein R₁₇ is $-C(O)-S-n\text{-octyl}$, and of formula 2.13, wherein Y₁ is nitrogen, Y₂, Y₃ and Y₄ are methine, R₂₄ is dimethylaminocarbonyl and Y₅ is methine, or Y₁, Y₂, Y₃ and Y₄ are methine, R₂₄ is methoxycarbonyl and Y₅ is methine, or Y₁, Y₂, Y₃ and Y₄ are methine, R₂₄ is 3-trifluoropropyl and Y₅ is nitrogen, or Y₁ is nitrogen, Y₂, Y₃ and Y₄ are methine, R₂₄ is ethylsulphonyl and Y₅ is methine, or Y₁ is N-Me, Y₂ is nitrogen, Y₃ and Y₄ together are C-Cl, R₂₄ is methoxycarbonyl and Y₅ is methine, and of formula 2.9, and of formula 2.11, wherein R₂₂ is trifluoromethyl, and of formula 2.12, and of formula 2.16, and of formula 2.18, and of formula 2.19, and

b) to antagonise the herbicide, an antidotally effective amount of a safener of formula 3.1

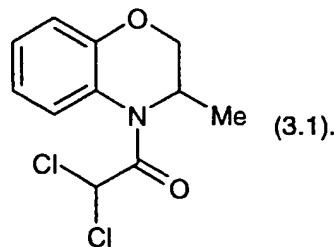


14. Herbicidal composition according to claim 9, which contains a) a herbicidally synergistic amount of a compound of formula I, a compound of formula 2.2



chloroacetyl-2-ethyl-6-methylaniline) and

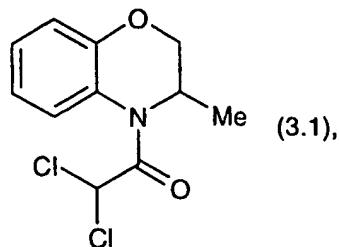
b) to antagonise the herbicide, an antidotally effective amount of a safener of formula 3.1



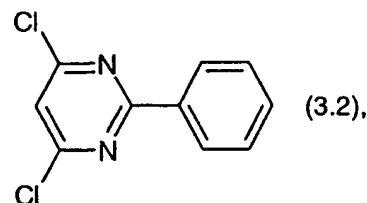
15. Selective herbicidal composition comprising, in addition to customary inert formulation assistants, such as carriers, solvents and wetting agents, as active ingredient a mixture of a) a herbicidally effective amount of a compound of formula I and

- 54 -

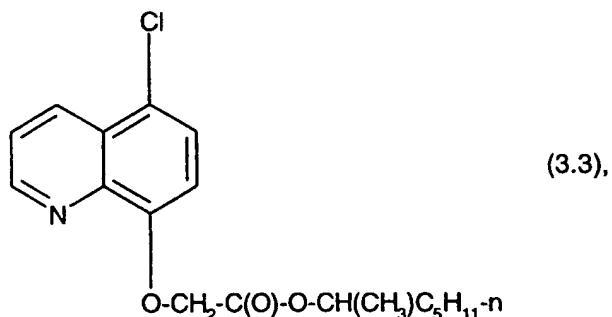
b) to antagonise the herbicide, an antidotally effective amount of a safener selected from the compound of formula 3.1



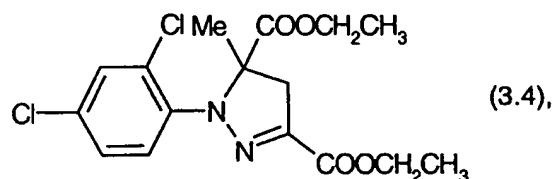
and the compound of formula 3.2



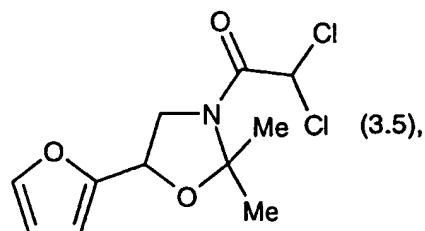
and the compound of formula 3.3



and the compound of formula 3.4

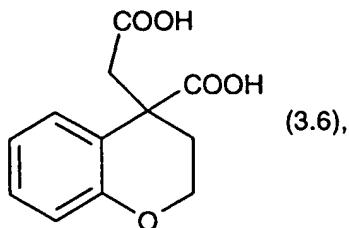


and the compound of formula 3.5

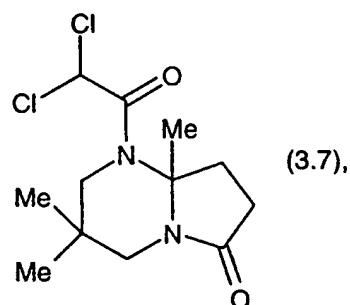


- 55 -

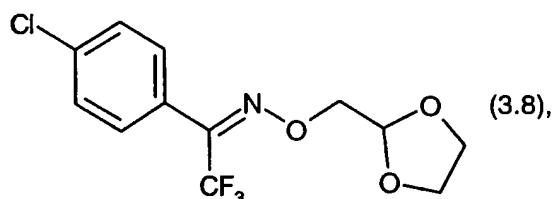
and the compound of formula 3.6



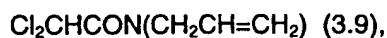
and the compound of formula 3.7



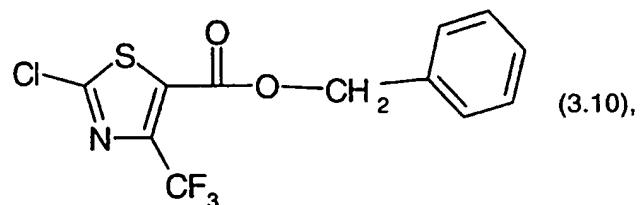
and the compound of formula 3.8



and of formula 3.9

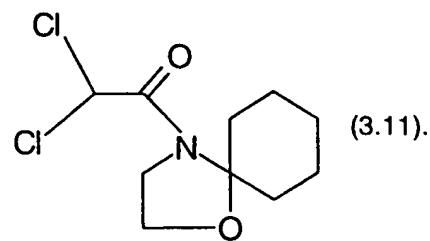


and of formula 3.10



and of formula 3.11

- 56 -



16. A method of selectively controlling weeds and grasses in crops of cultivated plants, which comprises treating said cultivated plants, the seeds or seedlings or the crop area thereof, with a herbicidally synergistic amount of a composition according to claim 15.

INTERNATIONAL SEARCH REPORT

International Application No

PCT/EP 99/04373

A. CLASSIFICATION OF SUBJECT MATTER

IPC 6 A01N41/10

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 6 A01N

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	EP 0 614 606 A (SANDOZ AG ;SANDOZ LTD (CH); SANDOZ AG (DE)) 14 September 1994 (1994-09-14) claims 1,5,9,11 ----	1,2,5-8
X	EP 0 298 680 A (ICI AMERICA INC) 11 January 1989 (1989-01-11) claims ----	15,16
X	EP 0 551 650 A (HOECHST AG) 21 July 1993 (1993-07-21) claims -----	15,16

 Further documents are listed in the continuation of box C. Patent family members are listed in annex.

* Special categories of cited documents :

- "A" document defining the general state of the art which is not considered to be of particular relevance
- "E" earlier document but published on or after the international filing date
- "L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)
- "O" document referring to an oral disclosure, use, exhibition or other means
- "P" document published prior to the international filing date but later than the priority date claimed

"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.

"Z" document member of the same patent family

Date of the actual completion of the international search

15 October 1999

Date of mailing of the international search report

27/10/1999

Name and mailing address of the ISA

European Patent Office, P.B. 5818 Patenttaan 2
NL - 2280 HV Rijswijk
Tel. (+31-70) 340-2040, Tx. 31 651 epo nl.
Fax: (+31-70) 340-3016

Authorized officer

Decorte, D

INTERNATIONAL SEARCH REPORT

Information on patent family members

Int'l Application No

PCT/EP 99/04373

Patent document cited in search report	Publication date	Patent family member(s)		Publication date
EP 0614606	A 14-09-1994	AU 672264	B	26-09-1996
		AU 5515794	A	25-08-1994
		BE 1007930	A	21-11-1995
		BR 9400602	A	23-08-1994
		BR 9402537	A	14-03-1995
		CA 2115863	A	19-08-1994
		CA 2126602	A	26-12-1994
		CN 1091240	A	31-08-1994
		CZ 9400345	A	14-09-1994
		DE 4421342	A	05-01-1995
		FR 2706735	A	30-12-1994
		FR 2727291	A	31-05-1996
		GB 2279872	A,B	18-01-1995
		HR 940114	A	31-10-1996
		HU 66739	A,B	28-12-1994
		HU 67562	A	28-04-1995
		IT 1272274	B	16-06-1997
		JP 7017807	A	20-01-1995
		MX 9404758	A	31-01-1995
		PL 302265	A	22-08-1994
		PL 176082	B	30-04-1999
		SG 49649	A	15-06-1998
		TR 960159	A	21-06-1996
		TR 960160	A	21-06-1996
		TR 960161	A	21-06-1996
		TR 960162	A	21-06-1996
		TR 960189	A	21-06-1996
		US 5877115	A	02-03-1999
		US 5491124	A	13-02-1996
		US 5545607	A	13-08-1996
		US 5922643	A	13-07-1999
		US 5888936	A	30-03-1999
		US 5908809	A	01-06-1999
		US 5895774	A	20-04-1999
		US 5888935	A	30-03-1999
		US 5905059	A	18-05-1999
		US 5905060	A	18-05-1999
		US 5928996	A	27-07-1999
		US 5900388	A	04-05-1999
		US 5900389	A	04-05-1999
		US 5885933	A	23-03-1999
		ZA 9404565	A	27-12-1995
		SK 18894	A	07-09-1994
		US 5716901	A	10-02-1998
		US 5721191	A	24-02-1998
		US 5895773	A	20-04-1999
		ZA 9401139	A	18-08-1995
EP 0298680	A 11-01-1989	US 4938796	A	03-07-1990
		AT 94339	T	15-10-1993
		AU 604336	B	13-12-1990
		AU 1867088	A	19-01-1989
		BG 60301	B	27-05-1994
		CA 1337158	A	03-10-1995
		CN 1034300	A,B	02-08-1989
		DE 3884076	D	21-10-1993
		DE 3884076	T	10-02-1994
		DK 376488	A	07-01-1989

INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No

PCT/EP 99/04373

Patent document cited in search report	Publication date	Patent family member(s)		Publication date
EP 0298680	A	EG	18668 A	30-12-1993
		ES	2059521 T	16-11-1994
		IE	61677 B	16-11-1994
		IL	86996 A	12-04-1994
		JP	1117803 A	10-05-1989
		JP	2896146 B	31-05-1999
		KR	9616186 B	06-12-1996
		MX	168198 B	11-05-1993
		NZ	225297 A	26-02-1991
		PH	25483 A	24-07-1991
		PT	87912 A, B	30-06-1989
		TR	24112 A	22-03-1990
		ZW	8988 A	14-03-1990
<hr/>		<hr/>		
EP 0551650	A	21-07-1993	AU	662581 B
			AU	3047792 A
			CA	2086491 A
			EP	0943240 A
			JP	5279204 A
			US	5441922 A
				07-09-1995
				08-07-1993
				01-07-1993
				22-09-1999
				26-10-1993
				15-08-1995